

DESCRIPTION

Melanin Concentrating Hormone Antagonist

5 TECHNICAL FIELD

The present invention relates to a melanin-concentrating hormone antagonist which is useful as an agent for preventing or treating obesity, etc.

10 BACKGROUND ART

Feeding behavior is an essential action for many living beings including humans. Therefore, if irregularities in feeding behavior occur, disorders, often connected to diseases, will occur in normal life-maintaining activities. Accompanying recent changes of our dietary environment, obesity is now becoming a social problem. In addition, not only is obesity a serious risk factor for life-style diseases such as diabetes, hypertension, and arteriosclerosis; it is also widely known that increased body weight places excessive burdens on joints such as knee joints, causing arthritis and pain.

The "diet boom," etc. show that there is a potentially great percentage of the population hoping to reduce body weight; on the other hand, many cases of feeding problems such as overeating, occurring due to causes such as hereditary neurosis or neurosis due to stress, have been reported.

Therefore, research on and development of agents for preventing or treating obesity, or agents for inhibiting eating, have been vigorously done for a long time.

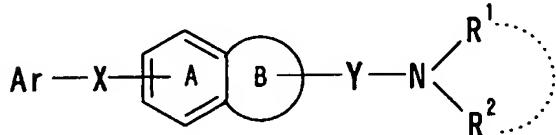
The centrally acting anorectic drug, Mazindol, is now being marketed.

Many appetite control factors such as leptin, have recently been discovered, and the development of anti-obesity agents or anorectic agents which will regulate the functions of these appetite control factors is progressing.

In particular, it is known that melanin-concentrating hormone (hereinafter also abbreviated as "MCH") originates in the hypothalamus and has orexigenic action. In addition, it has been reported that even though the daily behavior of MCH knock-out mice was normal, the amount of feeding by MCH knock-out mice was significantly reduced and their body weights were lighter than those of normal mice [Nature, Vol. 396, p.670, 1998]. This indicates that, if a MCH antagonist was produced, it can be expected to be an excellent anorectic agent or anti-obesity agent; but at present there are no known compound, especially non-peptide type compounds, which possess MCH antagonistic actions.

On the other hand, the following compounds are known as amine derivatives.

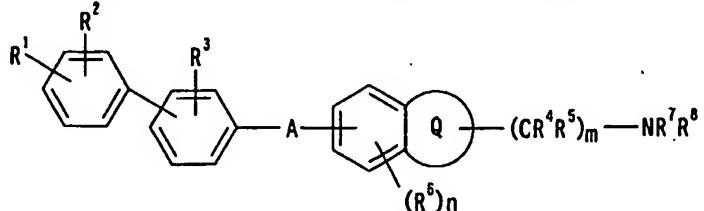
1) W098/38156 describes a compound of the formula :



wherein Ar is an optionally substituted ring assembly aromatic group or an optionally substituted condensed aromatic group; X is a bond, etc.; Y is an optionally substituted bivalent C₁₋₆ aliphatic hydrocarbon group which may have an intervening oxygen atom or sulfur atom; R¹ and R² are independently hydrogen atom or a lower alkyl, or R¹ and R², together with the adjacent nitrogen atom, form an optionally substituted nitrogen-containing hetero ring; Ring A is a benzene ring which may have further substituents in addition to the groups of the formula : -X-Ar where each symbol has the same meaning as defined above; Ring B is a 4 to 8 membered ring which may have further substituents in addition to the group of the formula : -Y-NR¹R² where each symbol has the same meaning as defined above; with the proviso that the condensed ring formed by ring A and ring B is an indole ring, the group of the formula : -X-Ar where

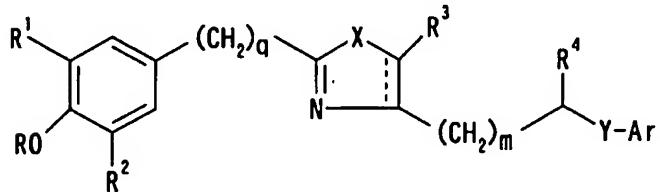
each symbol has the same meaning as defined above is substituted at the 4-, 6-, or 7- position on the indole ring; or its salt, which has an action of inhibiting the production and secretion of β -amyloid protein.

5 2) W095/32967 describes compound of the formula :



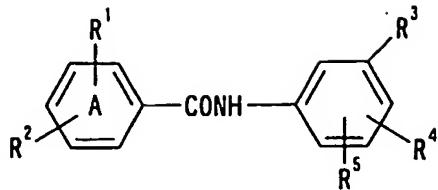
wherein A is CONR, in which R is hydrogen or C₁₋₆ alkyl; Q is an optionally substituted 5 to 7 membered hetero ring containing 1 to 3 hetero atoms selected from nitrogen or 10 sulfur; R¹ is hydrogen, halogen, etc.; R² and R³ are independently hydrogen, halogen, etc.; R₄ and R₅ are independently hydrogen or C₁₋₆ alkyl; R⁶ is halogen, hydroxy, etc.; R₇ and R₈ are independently hydrogen, C₁₋₆ alkyls, etc.; m is 0 to 4; n is 0, 1 or 2; or its salt, which has 5HT1D 15 antagonist activity and can be expected to ameliorate anorexia.

3) W098/15274 describes a compound of the formula :



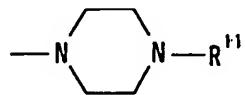
wherein Ar is phenyl, etc.; X is -O- or -S-; Y is CR⁵R^{5'}- 20 where R⁵' is H and R⁵ is -H, etc.; Z is -CH₂- or -N-; R is H or -(C_{1-C6}) alkyl; R¹ and R² are independently -(C_{1-C6}) alkyl, etc.; R³ is H etc.; R⁴ is hydrogen, etc.; m is an integer of 0 to 2; q is 0 or 1; n is an integer of 0 to 4; p is an integer of 1 to 6; t is an integer of 1 to 4; which 25 has an anti-oxidant activity and can be expected to ameliorate Alzheimer's disease.

4) EP533266



wherein R^1 is halogen, etc.; R^2 is phenyl optionally substituted by 1 or 2 substituents selected from halogen, etc.; R^3 is

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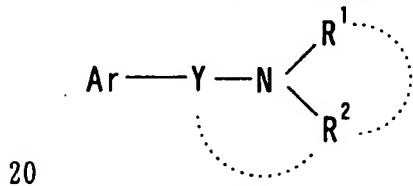


; R^4 and R^5 are independently hydrogen, halogen, etc.; R^{11} is hydrogen or C_{1-6} alkyl; which has 5HT1D antagonist activity, and can be expected to ameliorate anorexia.

There has been great desire for the development of a 10 melanin-concentrating hormone antagonist which is useful as an agent for preventing or treating obesity, excellent in oral absorbency, and safe.

DISCLOSURE OF INVENTION

15 As a result of intensive studies of compounds with a MCH antagonistic action, the present inventors found that a derivative which is obtained by introducing a group of the formula : $\text{Ar}^1-\text{X}-$ where each symbol has the same meaning as defined hereafter, into a compound of the formula :

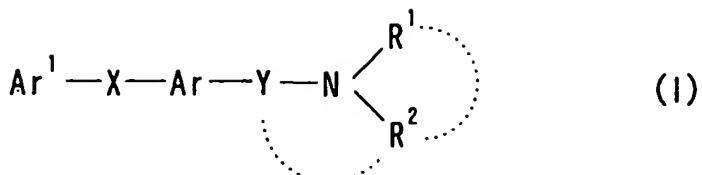


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wherein each symbol has the same meaning as defined hereinafter, had an excellent MCH antagonistic actions, to complete this invention.

Namely, the present invention relates to :

25 (1) a melanin-concentrating hormone antagonist which comprises a compound of the formula :



wherein Ar¹ is a cyclic group which may have substituents; X is a spacer having a main chain of 1 to 6 atoms; Y is a bond or a spacer having a main chain of 1 to 6 atoms;

5 Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have further substituents;

R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with 10 the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; R² may form a spiro ring together with Ar; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof;

15 (2) an antagonist according to the above (1), wherein Y is a spacer having a main chain of 1 to 6 atoms; R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring 20 which may have substituents; or R² may form a spiro ring together with Ar;

(3) an antagonist according to the above (2), wherein Ar¹ is an aromatic group which may have substituents; and "a hydrocarbon group which may have substituents" for R¹ and 25 R² is "C₁₋₆ alkyl which may have substituents";

(4) an antagonist according to the above (1), wherein the cyclic group for Ar¹ is C₆₋₁₄ monocyclic or condensed polycyclic aromatic hydrocarbon group;

(5) an antagonist according to the above (1), wherein the 30 cyclic group for Ar¹ is a group formed by removing an optional one hydrogen atom from an aromatic ring assemble in which 2 or 3 C₆₋₁₄ monocyclic or condensed polycyclic aromatic hydrocarbon groups are directly bonded by single

bonds;

(6) an antagonist according to the above (1), wherein the cyclic group for Ar¹ is a group formed by removing an optional one hydrogen atom from an aromatic ring assemble
5 in which C₆₋₁₄ monocyclic or condensed polycyclic aromatic hydrocarbon and 5 to 10 membered aromatic hetero ring are directly bonded by a single bond;

(7) an antagonist according to the above (1), wherein Ar¹ is phenyl, biphenylyl, phenyl-pyridyl, phenyl-furyl,
10 phenyl-isoxazolyl, diphenyl-oxazolyl, pyridyl-phenyl, phenyl-pyrimidinyl, benzofuranyl-phenyl, furyl-phenyl, terphenyl, thieryl-phenyl, indolyl, naphthyl-oxadiazolyl, benzofuranyl-oxadiazolyl, benzothienyl, benzofuranyl, fluorenyl, pyridyl-pyrrolyl or
15 thioxanthenyl;
each of which may have 1 to 3 substituents selected from the group consisting of halogen atom; nitro; C₁₋₃ alkylenedioxy; optionally halogenated C₁₋₆ alkyl; hydroxy-C₁₋₆ alkyl; optionally halogenated C₃₋₆ cycloalkyl;
20 optionally halogenated C₁₋₆ alkoxy; optionally halogenated C₁₋₆ alkythio; hydroxy; C₇₋₁₉ aralkyloxy which may have substituents; C₆₋₁₄ aryloxy which may have substituents; amino; mono-C₁₋₆ alkylamino; di-C₁₋₆ alkylamino; 5 to 7 membered saturated cyclic amino which may have substituents
25 and may be condensed with a benzene ring; 5 to 7 membered non-aromatic heterocyclic groups which may have substituents; formyl; carboxy; C₆₋₁₄ aryl-carbonyl which may have substituents; C₆₋₁₄ aryl-carbamoyl which may have substituents; aromatic hetero ring-carbamoyl which may have substituents; C₁₋₆ alkoxy-carbonyl; optionally halogenated C₁₋₆ alkyl-carboxamide; C₆₋₁₄ aryl-carboxamide which may have substituents; C₇₋₁₉ aralkyl-carboxamide which may have substituents; aromatic hetero ring-carboxamide which may have substituents; N-(C₆₋₁₄ aryl-carbonyl which
30 may have substituents)-N-C₁₋₆ alkylamino; C₆₋₁₄ arylamino-carbonylamino which may have substituents; C₆₋₁₄

arylsulfonylamino which may have substituents; C₆₋₁₄

aryl-carbonyloxy which may have substituents; oxo;

carboxy-C₁₋₆ alkyl; C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl; C₇₋₁₉

aralkyl which may have substituents; aromatic hetero

5 ring-C₁₋₆ alkoxy; and cyano;

(8) an antagonist according to the above (1), wherein Ar¹ is piperidinyl, piperazinyl, pyrrolidinyl, dihydropyridyl or tetrahydropyridyl; each of which may have 1 or 2 substituents selected from the group consisting of oxo, C₆₋₁₄

10 aryl which may have substituents, hydroxy, C₇₋₁₉, aralkyloxy-carbonyl, and C₇₋₁₉ aralkyl;

(9) an antagonist according to the above (1), wherein the "spacer having a main chain of 1 to 6 atoms" for X and Y is a bivalent group consisting of 1 to 3 species selected

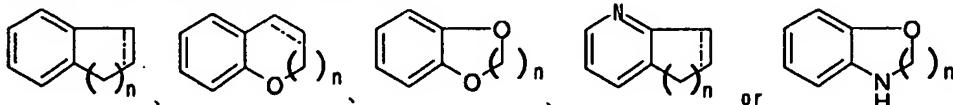
15 from -O-, -S-, -CO-, -SO-, -SO₂-, -NR⁸- (R⁸ is hydrogen atom, optionally halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkyl-carbonyl, optionally halogenated C₁₋₆ alkylsulfonyl), and a bivalent C₁₋₆ non-cyclic hydrocarbon group which may have substituents;

20 (10) an antagonist according to the above (1), wherein X is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}- wherein R^{8c} is hydrogen atom or C₁₋₆ alkyl;

(11) an antagonist according to the above (1), wherein Y is an optionally halogenated bivalent C₁₋₆ non-cyclic

25 hydrocarbon group;

(12) an antagonist according to the above (1), wherein Ar is a ring of the formula :

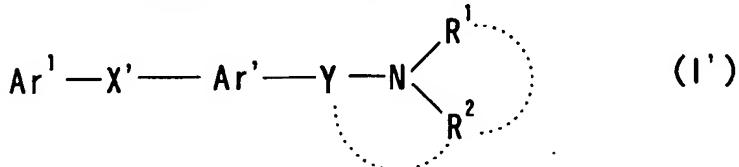


wherein _____ is a single bond or double bond, n is an integer of 1 to 4;

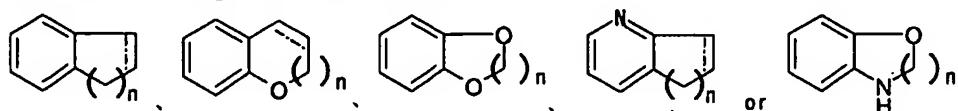
30 (13) an antagonist according to the above (1), wherein R¹ and R² are hydrogen atom or C₁₋₆ alkyl which may have substituents; or R¹ and R², together with the adjacent nitrogen atom, form a 3 to 8 membered nitrogen-containing

hetero ring;

- (14) an antagonist according to the above (1), which is an agent for preventing or treating diseases caused by a melanin-concentrating hormone;
- 5 (15) an antagonist according to the above (1), which is an agent for preventing or treating obesity;
- (16) an antagonist according to the above (1), which is an anorectic agent;
- (17) a pharmaceutical, which comprises a melanin-
- 10 concentrating hormone antagonist in combination with at least one species selected from the group consisting of an agent for treating diabetes, an agent for treating hypertension and an agent for treating arteriosclerosis;
- (18) a compound of the formula :



15 wherein Ar^1 is a cyclic group which may have substituents; Ar' is a ring of the formula :



wherein --- is a single bond or double bond, n is an integer of 1 to 4, and each ring may have substituents;

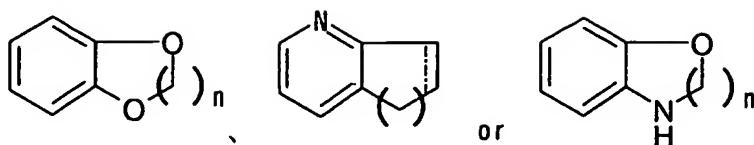
20 X' is $-\text{CONR}^{8c}-$, $-\text{NR}^{8c}\text{CO}-$, $-\text{CH}=\text{CH}-\text{CONR}^{8c}-$ or $-\text{SO}_2\text{NR}^{8c}-$ where R^{8c} is hydrogen atom or C_{1-6} alkyl;

Y is a spacer having a main chain of 1 to 6 atoms;

25 R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y , may form a nitrogen-containing hetero ring which may have

30 substituents;

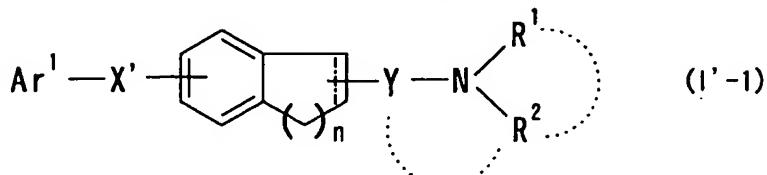
provided that Ar' is a ring of the formula :



wherein symbols have the same meanings as defined above, and each ring may have substituents, when X' is $\text{-SO}_2\text{NH-}$; and provided that Ar¹ is not biphenylyl which may be

5 substituted, when X' is -CONH- and Ar' is any one of benzopyran, dihydrobenzopyran, dihyrobenzoxazine, dihydrobenzoxazole or tetrahydrobenzoxazepine; (excluding N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-4-biphenylylcarboxamide); or a salt thereof;

10 (19) a compound of the formula :



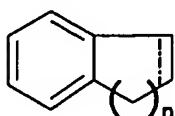
wherein Ar¹ is a cyclic group which may have substituents;

----- is a single bond or double bond;

n is an integer of 1 to 4;

15 X' is $\text{-CONR}^{8c}\text{-}$, $\text{-NR}^{8c}\text{CO-}$ or $\text{-CH=CH-CNR}^{8c}\text{-}$ where R^{8c} is hydrogen atom or C₁₋₆ alkyl; Y is a spacer having a main chain of 1 to 6 atoms; R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;

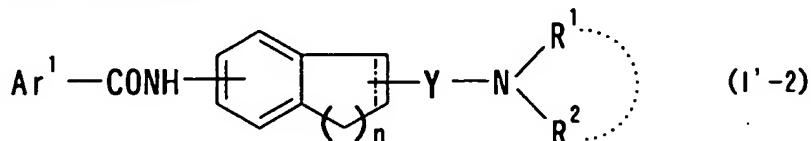
20 25 a ring of the formula :



wherein symbols have the same meanings as defined above,

may have further substituents;
 provided that N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-4-biphenylylcarboxamide is excluded; or a salt thereof;

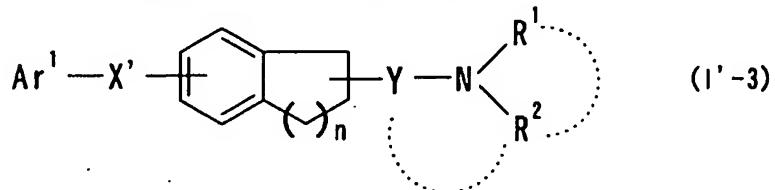
5 (20) a compound according to the above (19), which is of the formula :



wherein R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; the other symbols have the same meanings as defined in the above (19);

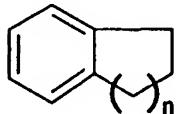
10 (21) a compound according to the above (20), wherein Ar¹ is an aromatic group which may have substituents; and "a hydrocarbon group which may have substituents" for R¹ and R² is "C₁₋₆ alkyl which may have substituents";

15 (22) a compound of the formula :



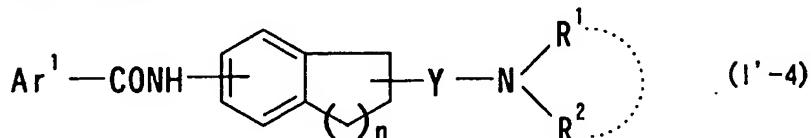
20 wherein Ar¹ is a cyclic group which may have substituents; n is an integer of 1 to 4; X' is -CONR^{8c}-, -NR^{8c}CO- or -CH=CH-CO-NR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl; Y is a spacer having a main chain of 1 to 6 atoms; 25 R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R², together with the adjacent nitrogen atom and Y, may form a 30 nitrogen-containing hetero ring which may have

substituents;
a ring of the formula :



wherein n has the same meaning as defined above, may have
5 further substituents;
provided that N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-4-biphenylcarboxamide is excluded; or a salt thereof;

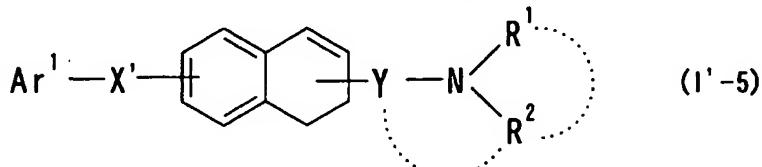
(23) a compound according to the above (22), which is of
10 the formula :



wherein R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a
15 nitrogen-containing hetero ring which may have substituents; the other symbols have the same meanings as defined in the above (22);

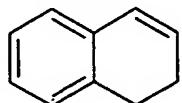
(24) a compound according to the above (23), wherein Ar¹ is an aromatic group which may have substituents; and "a hydrocarbon group which may have substituents" for R¹ and R² is "C₁₋₆ alkyl which may have substituents";

(25) a compound of the formula :

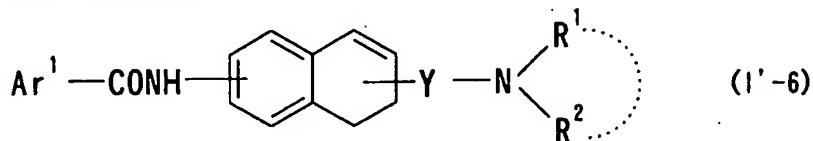


wherein Ar¹ is a cyclic group which may have substituents;
25 X' is -CONR^{8c}-, -NR^{8c}CO- or -CH=CH-CONR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl;
Y is a spacer having a main chain of 1 to 6 atoms;
R¹ and R² are independently hydrogen atom or a hydrocarbon

group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R², together with the adjacent nitrogen atom and Y, may form a
 5 nitrogen-containing hetero ring which may have substituents;
 a ring of the formula :



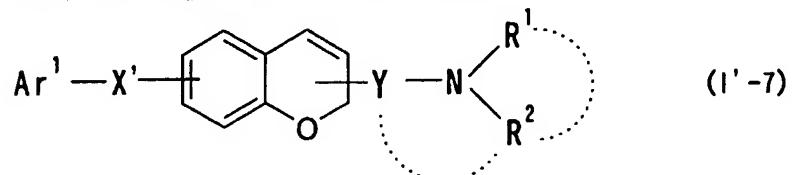
may have further substituents; or a salt thereof;
 10 (26) a compound according to the above (25), which is of the formula :



wherein R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; the other symbols have the same meanings as defined in the above (25);

(27) a compound according to the above (26), wherein Ar¹ is an aromatic group which may have substituents; and "a hydrocarbon group which may have substituents" for R¹ and R² is "C₁₋₆ alkyl which may have substituents";

(28) a compound of the formula :

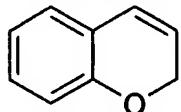


25 wherein Ar¹ is a cyclic group which may have substituents; X' is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl;
 Y is a spacer having a main chain of 1 to 6 atoms;

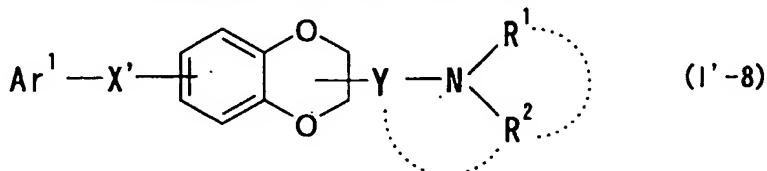
R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;

5 with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;

a ring of the formula :



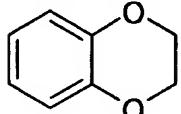
10 may have further substituents;
provided that Ar^1 is not biphenylyl which may be substituted, when X' is -CONH-; or a salt thereof;
(29) a compound of the formula :



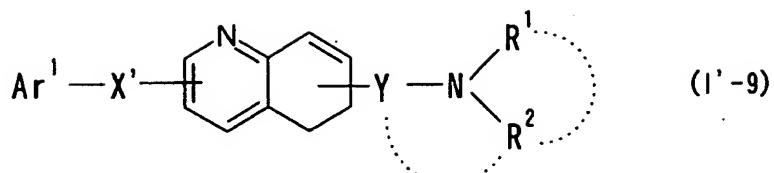
15 wherein Ar^1 is a cyclic group which may have substituents;
X' is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CO NR^{8c}- or -SO₂NR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl;
Y is a spacer having a main chain of 1 to 6 atoms;
R¹ and R² are independently hydrogen atom or a hydrocarbon
20 group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;

25 with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;

a ring of the formula :



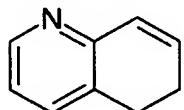
may have further substituents; or a salt thereof;
(30) a compound of the formula :



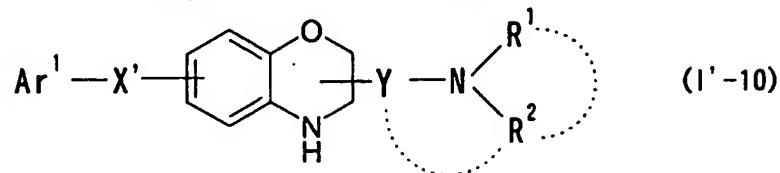
wherein Ar¹ is a cyclic group which may have substituents; X' is -CONR^{8c}- , -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl;

- 5 Y is a spacer having a main chain of 1 to 6 atoms; R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R², together
- 10 with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;

a ring of the formula :



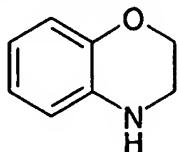
- 15 may have further substituents; or a salt thereof:
- (31) a compound of the formula :



wherein Ar¹ is a cyclic group which may have substituents; X' is -CONR^{8c}- , -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl;

- Y is a spacer having a main chain of 1 to 6 atoms; R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R², together
- 20 with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;
- 25

a ring of the formula :



may have further substituents;

provided that Ar¹ is not biphenyl which may be

5 substituted, when X' is -CONH-; or a salt thereof;

(32) a pharmaceutical composition which comprises a compound as defined in any one of the above (18), (19), (22), (25), (26), (28), (29), (30) and (31);

10 (33) a prodrug of a compound as defined in any one of the above (18), (19), (22), (25), (26), (28), (29), (30) and (31);

(34) a compound according to the above (18), which is N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-(4'-methoxybiphenyl-4-yl)carboxamide;

15 4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;

4'-fluoro-N-[6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]4-carboxamide;

4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-

20 tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;

(+)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;

25 (-)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;

4'-chloro-N-[3-[(N,N-dimethylamino)methyl]-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide;

30 4'-fluoro-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;

N-[3-[(dimethylamino)methyl]-2H-chromen-7-yl]-4'-fluoro[1,1'-biphenyl]-4-carboxamide;

4'-chloro-N-[6-[(dimethylamino)methyl]-5-methyl-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
6-(4-methoxyphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide;
5 naphthalenyl]nicotinamide;
4'-chloro-N-[7-[(dimethylamino)methyl]-5,6-dihydro-3-quinolinyl][1,1'-biphenyl]-4-carboxamide;
4-(4-chlorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-3,6-dihydro-1(2H)-
10 pyridinecarboxamide;
N-[6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]-4-(4-fluorophenyl)-1-piperidinecarboxamide;
4-(4-methoxyphenyl)-N-[6-(1-pyrrolidinylmethyl)-5-methyl-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide;
15 4'-fluoro-N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
4'-chloro-N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
20 4'-chloro-N-[2-[(dimethylamino)methyl]-3,4-dihydro-2H-1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide;
4-(4-methoxyphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide;
25 4-(4-chlorophenyl)-N-[6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide;
4'-chloro-N-[2-[(dimethylamino)methyl]-1H-inden-6-yl][1,1'-biphenyl]-4-carboxamide;
30 4'-fluoro-N-[2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide;
4'-fluoro-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
35 4'-chloro-N-[5-methyl-6-[(4-methyl-1-

5 4-(4-chlorophenyl)-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide;

10 (35) a method for preventing or treating diseases caused by a melanin-concentrating hormone in a mammal in need thereof, which comprises administering to said mammal an effective amount of a compound or a salt thereof as defined in the above (1);

15 (36) a method for preventing or treating obesity in a mammal in need thereof, which comprises administering to said mammal an effective amount of a compound or a salt thereof as defined in the above (1);

20 (37) use of a compound or a salt thereof as defined in the above (1), for the manufacture of a pharmaceutical preparation for preventing or treating diseases caused by a melanin-concentrating hormone; and

25 (38) use of a compound or a salt thereof as defined in the above (1), for the manufacture of a pharmaceutical preparation for preventing or treating obesity.

25 Examples of "cyclic group" in the "cyclic group which may have substituents" for Ar¹ include aromatic groups, non-aromatic cyclic hydrocarbon groups, non-aromatic heterocyclic groups.

30 Here, examples of "aromatic groups" include monocyclic aromatic groups, condensed aromatic groups, and ring assembly aromatic groups.

35 Examples of the concerned monocyclic aromatic groups include univalent groups which can be formed by removing an optional one hydrogen atom from a monocyclic aromatic ring. Example of the "monocyclic aromatic ring" include a benzene ring and a 5 or 6 membered aromatic hetero ring.

40 Examples of the "5 or 6 membered aromatic hetero ring" include a 5 or 6 membered aromatic hetero ring containing

one or more (for example, 1 to 3) hetero atom selected from nitrogen, sulfur and oxygen atom in addition to a carbon atom. Concretely, thiophene, furan, pyrrole, imidazole, pyrazole, thiazole, isothiazole, oxazole, isoxazole, 5 pyridine, pyrazine, pyrimidine, pyridazine, 1,2,4-oxadiazole, 1,3,4-oxadiazole, 1,2,4-thiadiazole, 1,3,4-thiadiazole, furazan, etc., can be mentioned.

Concrete examples of the "monocyclic aromatic groups" include phenyl, 2- or 3-thienyl, 2-, 3-, or 4-pyridyl, 2-10 or 3-furyl, 2-, 4- or 5-thiazonyl, 2-, 4- or 5-oxazolyl, 1-, 3- or 4-pyrazolyl, 2-pyrazinyl, 2-, 4- or 5-pyrimidinyl, 1-, 2- or 3-pyrrolyl, 1-, 2- or 4-imidazolyl, 3- or 4-pyridazinyl, 3-isothiazolyl, 3-isooxazolyl, 1,2,4-oxadiazol-5-yl, 1,2,4-oxadiazol-3-yl.

15 The "condensed aromatic groups" mean a univalent group that can be formed by removing an optional one hydrogen atom from condensed polycyclic (preferably bicyclic to tetracyclic, more preferably bicyclic or tricyclic) aromatic rings. Examples of the "condensed aromatic 20 groups" include condensed polycyclic aromatic hydrocarbons, condensed polycyclic aromatic hetero rings.

Examples of the "condensed polycyclic aromatic hydrocarbons" include C₉₋₁₄ condensed polycyclic (bicyclic or tricyclic) aromatic hydrocarbons (e.g. naphthalene, 25 indene, fluorene, anthracene, etc.).

Examples of the "condensed polycyclic aromatic hetero rings" include 9 to 14 membered, preferably, 9 or 10 membered, condensed polycyclic aromatic hetero rings containing one or more (for instance, 1 to 4 atoms) hetero 30 atoms selected from nitrogen, sulfur and oxygen atom in addition to carbon atoms. Concrete examples of the "condensed polycyclic aromatic hetero rings" include benzofuran, benzimidazole, benzoxazole, benzothiazole, benzisothiazole, naphtho[2,3-b]thiophene, isoquinoline, 35 quinoline, indole, quinoxaline, phenanthridine, phenothiadine, phenoxazine, phthaladine, naphthylidine,

quinazoline, cinnoline, carbazole, β -carboline, acridine, phenazine, phthalimide, thioxanthene.

Concrete examples of "condensed aromatic groups" include 1-naphthyl; 2-naphthyl; 2-, 3-, 4-, 5- or 8-5 quinolyl; 1-, 3-, 4-, 5-, 6-, 7- or 8-isoquinolyl; 1-, 2-, 3-, 4-, 5-, 6- or 7-indolyl; 1-, 2-, 4- or 5-isoindolyl; 1-, 5- or 6-phthalazinyl; 2-, 3- or 5-quinoxalinyl; 2-, 3-, 4-, 5- or 6-benzofuranyl; 2-, 4-, 5- or 6-benzothiazolyl; 1-, 2-, 4-, 5- or 6-benzimidazolyl; 1-, 2-, 3- or 4-10 fluorenyl; thioxanthenyl.

"Ring assembly aromatic group" means a group formed by removing an optional one hydrogen atom from an aromatic ring assembles in which 2 or more (preferably 2 or 3) aromatic rings are directly bonded by single bonds, and in 15 which the number of bonds which directly bond the rings, is less by one than the number of ring systems.

Examples of the aromatic ring assembles include an aromatic ring assembles formed by 2 or 3 (preferably 2) species selected from C₆₋₁₄ monocyclic or condensed 20 polycyclic aromatic hydrocarbons (e.g. benzene and naphthalene) and 5 to 10 membered (preferably 5 or 6 membered) aromatic hetero rings.

Preferable example of the aromatic ring assembles include aromatic ring assembles comprising 2 or 3 aromatic 25 rings selected from benzene, naphthalene, pyridine, pyrimidine, thiophene, furan, thiazole, isothiazole, oxazole, isoxazole, 1,2,4-oxadiazole, 1,3,4-oxadiazole, 1,2,4-thiadiazole, 1,3,4-thiadiazole, quinoline, isoquinoline, indole, benzothiophene, benzoxazole, 30 benzothiazole, benzofuran and pyrrole.

Concrete examples of the "ring assembly aromatic groups" include 2-, 3- or 4-biphenyl; 3-(1-naphthyl)-1,2,4-oxadiazol-5-yl; 3-(2-naphthyl)-1, 2, 4-oxadiazol-5-yl; 3-(2-benzofuranyl)-1,2,4-oxadiazol-5-yl; 3-35 phenyl-1,2,4-oxadiazol-5-yl; 3-(2-benzoxazolyl)-1,2,4-oxadiazol-5-yl; 3-(3-indolyl)-1,2,4-oxadiazol-5-yl; 3-

(2-indolyl)-1,2,4-oxadiazol-5-yl; 4-phenylthiazol-2-yl; 4-(2-benzofuranyl)thiazol-2-yl; 4-phenyl-1,3-oxazol-5-yl; 5-phenyl-isothiazol-4-yl; 5-phenyloxazol-2-yl; 4-(2-thienyl)phenyl; 4-(3-thienyl)phenyl; 3-(3-pyridyl)phenyl; 4-(3-pyridyl)phenyl; 6-phenyl-3-pyridyl; 5-phenyl-1,3,4-oxadiazol-2-yl; 4-(2-naphthyl)phenyl; 4-(2-benzofuranyl)phenyl; 4,4'-terphenyl; 5-phenyl-2-pyridyl; 2-phenyl-5-pyrimidinyl; 4-(4-pyridyl)phenyl; 2-phenyl-1,3-oxazol-5-yl; 2,4-diphenyl-1,3-oxazol-5-yl; 10 3-phenyl-isoxazol-5-yl; 5-phenyl-2-furyl; 4-(2-furyl)phenyl; 3-(4-pyridyl)pyrrolyl.

Preferable groups among the above "aromatic groups" are " C_{6-14} monocyclic or condensed polycyclic aromatic hydrocarbon groups (preferably, phenyl, etc.)", "a group formed by removing an optional one hydrogen atom from an aromatic ring assembles in which 2 or 3 C_{6-14} monocyclic or condensed polycyclic aromatic hydrocarbon groups are directly bonded by single bonds (preferably, 2-, 3- or 4-biphenylyl; 4,4-terphenyl, etc.)" and "a group formed by removing an optional one hydrogen atom from an aromatic ring assembles in which a C_{6-14} monocyclic or condensed polycyclic aromatic hydrocarbon and 5 to 10 membered aromatic hetero ring are directly bonded by a single bond (preferably, 6-phenyl-3-pyridyl, 5-phenyl-2-pyridyl, etc.)".

25 Examples of "non-aromatic cyclic hydrocarbon groups" include C_{3-8} Cycloalkyl, C_{3-8} cycloalkenyl.

Here, concrete examples of C_{3-8} cycloalkyl include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl.

30 Concrete examples of C_{3-8} cycloalkenyl include cyclopropenyl, cyclobutenyl, cyclopentenyl, cyclohexenyl, cycloheptenyl, cyclooctenyl.

Among the above "non-aromatic cyclic hydrocarbon groups", C_{3-8} cycloalkyl is preferable, and cyclohexyl is particularly preferable.

35 Examples of "non-aromatic heterocyclic groups"

include monocyclic non-aromatic heterocyclic groups, condensed polycyclic non-aromatic heterocyclic groups.

Examples of the "monocyclic non-aromatic heterocyclic groups" include univalent groups formed by removing an optional one hydrogen atom from monocyclic non-aromatic hetero ring. Examples of the "monocyclic non-aromatic heterocyclic groups" include 5 to 8 membered monocyclic non-aromatic heterocyclic groups containing one or more (e.g. 1 to 3) hetero atoms selected from nitrogen, sulfur and oxygen atom in addition to carbon atoms. Concretely, tetrahydrothiophene, tetrahydrofuran, pyrrolidine, imidazoline, imidazolidine, pyrazoline, pyrazolidine, tetrahydrothiazole, tetrahydroisothiazole, tetrohydrooxazole, tetrohydroisoxazole, piperidine, tetrahydropyridine, dihydropyridine, piperazine, morpholine, thiomorpholine, tetrahydropyrimidine, tetrahydropyridazine, hexamethyleneimine, etc. can be mentioned.

"Condensed polycyclic non-aromatic heterocyclic group" means a univalent group formed by removing an optional one hydrogen atom from a condensed polycyclic (preferably bicyclic to tetracyclic, more preferably bicyclic or tricyclic) non-aromatic hetero ring. Examples of the "condensed polycyclic non-aromatic hetero ring" include 9 to 14 membered, preferably 9 or 10 membered condensed polycyclic non-aromatic hetero rings which contain one or more (e.g. 1 to 4) hetero atoms selected from nitrogen, sulfur and oxygen atom in addition to carbon atoms.

Concretely, dihydrobenzofuran, dihydrobenzimidazole, dihydrobenzoxazole, dihydrobenzothiazole, dihydrobenzisothiazole, dihydronaphtho[2,3-b]thiophene, tetrahydroisoquinoline, tetrahydroquinoline, indoline, isoindoline, tetrahydroquinoxaline, tetrahydrophenanthridine, hexahydrophenothiadine, hexahydrophenoxazine,

tetrahydropthaladine, tetrahydronaphthylidine,
tetrahydroquinazoline, tetrahydrocinnoline,
tetrahydrocarbazole, tetrahydro- β -carboline,
tetrahydroacridine, tetrahydrophenazine,
5 tetrahydrothioxantene, etc., can be mentioned.
Among the above "non-aromatic heterocyclic groups",
"5 to 8 membered monocyclic non-aromatic heterocyclic
groups (preferably piperidinyl; piperazinyl;
pyrrolidinyl; dihydropyridyl; tetrahydropyridyl, etc.)"
10 are preferable.

Examples of "substituents" in the "cyclic group which
may have substituents" for Ar¹ include oxo, halogen atoms
(e.g. fluorine, chlorine, bromine, iodine, etc.), C₁₋₃,
15 alkylenedioxy (e.g. methylenedioxy, ethylenedioxy, etc.),
nitro, cyano, optionally halogenated C₁₋₆ alkyl, hydroxy-C₁₋₆
alkyl, carboxy-C₁₋₆ alkyl, C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl,
C₆₋₁₄ aryloxy-C₁₋₆ alkyl (e.g. phenoxyethyl, etc.), C₁₋₆
alkyl-C₆₋₁₄ aryl-C₂₋₆ alkenyl (e.g. methylphenylethenyl,
20 etc.), optionally halogenated C₃₋₆ cycloalkyl, optionally
halogenated C₁₋₆ alkoxy, optionally halogenated C₁₋₆
alkylthio, C₇₋₁₉ aralkyl which may have substituents,
hydroxy, C₆₋₁₄ aryloxy which may have substituents, C₇₋₁₉,
aralkyloxy which may have substituents, C₆₋₁₄ aryl-carbamoyl
25 which may have substituents, amino, amino-C₁₋₆ alkyl (e.g.
aminomethyl, aminoethyl, aminopropyl, aminobutyl, etc.).
mono-C₁₋₆ alkylamino (e.g. methylamino, ethylamino,
propylamino, isopropylamino, butylamino, etc.), di-C₁₋₆
alkylamino (e.g. dimethylamino, diethylamino,
30 dipropylamino, dibutylamino, ethylmethylamino, etc.),
mono-C₁₋₆ alkylamino-C₁₋₆ alkyl (e.g. methylaminomethyl,
ethylaminomethyl, propylaminomethyl,
isopropylaminoethyl, butylaminoethyl, etc.), di-C₁₋₆
alkylamino-C₁₋₆ alkyl (e.g. dimethylaminomethyl,
35 diethylaminomethyl, dipropylaminomethyl,
diisopropylaminoethyl, dibutylaminoethyl, etc.), 5 to 7

membered saturated cyclic amino which may have substituents, 5 to 7 membered non-aromatic heterocyclic groups which may have substituents, acyl, acylamino, acyloxy, aromatic hetero ring-C₁₋₆ alkoxy.

5 The "cyclic group" for Ar¹ may have 1 to 5, preferably 1 to 3, of the above-mentioned substituents at a substitutable position on the cyclic group. When the number of substituents is 2 or more, each substituents can be the same or different.

10 Also, when the "cyclic group" for Ar¹ is a non-aromatic cyclic hydrocarbon group or a non-aromatic heterocyclic group, the "cyclic group" may have as its substituents, C₆₋₁₄ aryl which may have substituents, and 5 to 10 membered aromatic heterocyclic groups which may have substituents.

15 Here, the groups exemplified as "substituents" in the "5 to 7 membered saturated cyclic amino which may have substituents" mentioned hereinafter, can be mentioned as "C₆₋₁₄ aryl which may have substituents" and "5 to 10 membered aromatic heterocyclic groups which may have substituents".

20 The number of substituents is, for instance, 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Concrete examples of the above "optionally halogenated C₁₋₆ alkyl" include C₁₋₆ alkyl (e.g. methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, hexyl, etc.) which may have 1 to 5, preferably 1 to 3, halogen atoms (e.g. fluorine, chlorine, bromine, iodine, etc.). Concrete examples include methyl, chloromethyl, difluoromethyl, trichloromethyl, trifluoromethyl, ethyl, 2-bromoethyl, 2,2,2-trifluoroethyl, pentafluoroethyl, propyl, 3,3,3-trifluoropropyl, isopropyl, butyl, 4,4,4-trifluorobutyl, isobutyl, sec-butyl, tert-butyl, pentyl, isopentyl, neopentyl, 5,5,5-trifluoropentyl, hexyl, 6,6,6-trifluorohexyl.

35 The C₁₋₆ alkyl in the above "optionally halogenated C₁₋₆

alkyl" can be mentioned as the C₁₋₆ alkyl in the above "hydroxy-C₁₋₆ alkyl", "carboxy-C₁₋₆ alkyl" and "C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl". Examples of C₁₋₆ alkoxy in the "C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl" include methoxy, ethoxy, 5 propoxy, butoxy, pentyloxy.

Examples of the above "optionally halogenated C₃₋₆ cycloalkyl" include C₃₋₆ cycloalkyl (e.g. cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, etc.) which may have 1 to 5, preferably 1 to 3, halogen atoms (e.g. fluorine, 10 chlorine, bromine, iodine, etc.). Concrete examples include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, 4,4-dichlorocyclohexyl, 2,2,3,3-tetrafluorocyclopentyl, 4-chlorocyclohexyl.

Examples of the above "optionally halogenated C₁₋₆ alkoxy" include C₁₋₆ alkoxy (e.g. methoxy, ethoxy, propoxy, butoxy, pentyloxy, etc.) which may have 1 to 5, preferably 1 to 3, halogen atoms (e.g. fluorine, chlorine, bromine, iodine, etc.). Concrete examples include methoxy, difluoromethoxy, trifluoromethoxy, ethoxy, 2,2,2-trifluoroethoxy, propoxy, isopropoxy, butoxy, 4,4,4-trifluorobutoxy, isobutoxy, sec-butoxy, pentyloxy, hexyloxy.

Examples of the above "optionally halogenated C₁₋₆ alkylthio" include C₁₋₆ alkylthio (e.g. methylthio, 25 ethylthio, propylthio, isopropylthio, butylthio, sec-butylthio, tert-butylthio, etc.) which may have 1 to 5, preferably 1 to 3, halogen atoms (e.g. fluorine, chlorine, bromine, iodine, etc.). Concrete examples include methylthio, difluoromethylthio, trifluoromethylthio, 30 ethylthio, propylthio, isopropylthio, butylthio, 4,4,4-trifluorobutylthio, pentylthio, hexylthio.

Examples of the "C₇₋₁₉, aralkyl" in the above "C₇₋₁₉ aralkyl which may have substituents" include benzyl, phenethyl, diphenylmethyl, triphenylmethyl, 1-naphthylmethyl, 2-naphthylmethyl, 2,2-diphenylethyl, 3-phenylpropyl, 4-phenylbutyl, 5-phenylpentyl. Benzyl is

particularly preferable.

Examples of the "substituents" in the above "C₇₋₁₉, aralkyl which may have substituents" include halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), C₁₋₃, 5 alkylene dioxy (e.g. methylenedioxy, ethylenedioxy, etc.), nitro, cyano, optionally halogenated C₁₋₆ alkyl, optionally halogenated C₃₋₆ cycloalkyl, optionally halogenated C₁₋₆ alkoxy, optionally halogenated C₁₋₆ alkylthio, hydroxy, amino, mono-C₁₋₆ alkylamino (e.g. methylamino, ethylamino, 10 propylamino, isopropylamino, butylamino, etc.), di-C₁₋₆ alkylamino (e.g. dimethylamino, diethylamino, dipropylamino, dibutylamino, ethylmethylamino, etc.), amino-C₁₋₆ alkyl (e.g. aminomethyl, aminoethyl, aminopropyl, aminobutyl, etc.), mono-C₁₋₆ alkylamino-C₁₋₆, 15 alkyl (e.g. methylaminomethyl, ethylaminomethyl, propylaminomethyl, isopropylaminoethyl, butylaminoethyl, etc.), di-C₁₋₆ alkylamino-C₁₋₆ alkyl (e.g. dimethylaminomethyl, diethylaminomethyl, dipropylaminomethyl, diisopropylaminoethyl, 20 dibutylaminoethyl, etc.), formyl, carboxy, carbamoyl, thiocabamoyl, optionally halogenated C₁₋₆ alkyl-carbonyl, C₁₋₆ alkoxy-carbonyl (e.g., methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, tert-butoxycarbonyl, etc.), mono-C₁₋₆ alkyl-carbamoyl (e.g., methylcarbamoyl, 25 ethylcarbamoyl, etc.), di-C₁₋₆ alkyl-carbamoyl (e.g. dimethylcarbamoyl, diethylcarbamoyl, ethylmethylecarbamoyl, etc.), optionally halogenated C₁₋₆ alkylsulfonyl, formylamino, optionally halogenated C₁₋₆ alkyl-carboxamide, C₁₋₆ alkoxy-carboxamide (e.g. 30 methoxycarboxamide, ethoxycarboxamide, propoxycarboxamide, butoxycarboxamide, etc.), C₁₋₆ alkylsulfonylamino (e.g. methylsulfonylamino, ethylsulfonylamino, etc.), C₁₋₆ alkyl-carbonyloxy (e.g. acetoxy, propanoyloxy, etc.), C₁₋₆ alkoxy-carbonyloxy (e.g. 35 methoxycarbonyloxy, ethoxycarbonyloxy, propoxycarbonyloxy, butoxycarbonyloxy, etc.) mono-C₁₋₆

alkyl-carbamoyloxy (e.g. methylcarbamoyloxy, ethylcarbamoyloxy, etc.), di-C₁₋₆ alkyl-carbamoyloxy (e.g. dimethylcarbamoyloxy, diethylcarbamoyloxy, etc.). The number of substituents is, for instance, 1 to 5, preferably 5 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

As "optionally halogenated C₁₋₆ alkyl", "optionally halogenated C₃₋₆ cycloalkyl", "optionally halogenated C₁₋₆ alkoxy" and "optionally halogenated C₁₋₆ alkylthio", those 10 exemplified as "substituents" in the above "cyclic group which may have substituents" can be used respectively.

Examples of the above "optionally halogenated C₁₋₆ alkylcarbonyl" include C₁₋₆ alkyl-carbonyl (e.g. acetyl, propanoyl, butanoyl, pentanoyl, hexanoyl, etc.) which may 15 have 1 to 5, preferably 1 to 3, halogen atoms (e.g., fluorine, chlorine, bromine, iodine, etc.). Concrete examples include acetyl, monochloroacetyl, trifluoroacetyl, trichloroacetyl, propanoyl, butanoyl, pentanoyl, hexanoyl.

20 Examples of the above "optionally halogenated C₁₋₆ alkylsulfonyl" include C₁₋₆ alkylsulfonyl (e.g. methysulfonyl, ethylsulfonyl, propylsulfonyl, isopropylsulfonyl, butylsulfonyl, sec-butylsulfonyl, tert-butylsulfonyl, etc.) which may have 1 to 5, preferably 25 1 to 3, halogen atoms (e.g., fluorine, chlorine, bromine, iodine, etc.). Concrete examples include methysulfonyl, difluoromethylsulfonyl, trifluoromethylsulfonyl, ethylsulfonyl, propylsulfonyl, isopropylsulfonyl, butylsulfonyl, 4,4,4-trifluorobutylsulfonyl, pentylsulfonyl, hexylsulfonyl.

30 Examples of the above "optionally halogenated C₁₋₆ alkyl-carboxamide" include C₁₋₆ alkyl-carboxamide (e.g. acetamide, propanamide, butanamide, etc.) which may have 1 to 5, preferably 1 to 3, halogen atoms (e.g. fluorine, 35 chlorine, bromine, iodine, etc.). Concrete examples include acetamide, trifluoroacetamide, propanamide,

butanamide.

Examples of "C₆₋₁₄ aryloxy" in the above "C₆₋₁₄ aryloxy which may have substituents" include phenoxy, 1-naphthoxy, 2-naphthoxy.

5 Examples of "C₇₋₁₉ aralkyloxy" in the above "C₇₋₁₉ aralkyloxy which may have substituents" include benzyloxy, phenethyloxy, diphenylmethyloxy, triphenylmethyloxy, 1-naphthylmethyloxy, 2-naphthylmethyloxy, 2,2-diphenylethyloxy, 3-phenylpropyloxy, 4-phenylbutyloxy,
10 5-phenylpentyloxy.

Examples of "C₆₋₁₄ arylcarbamoyl" in the above "C₆₋₁₄ arylcarbamoyl which may have substituents" include phenylcarbamoyl, 1-naphthylcarbamoyl, 2-naphthylcarbamoyl.

15 As the "substituents" in the "C₆₋₁₄ aryloxy which may have substituents", "C₇₋₁₉ aralkyloxy which may have substituents" and "C₆₋₁₄ aryl-carbamoyl which may have substituents", those exemplified for "substituents" in the above "C₇₋₁₉ aralkyl which may have substituents" can be used.

20 The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Examples of the "5 to 7 membered saturated cyclic amino" in the above "5 to 7 membered saturated cyclic amino which may have substituents" include morpholino, thiomorpholino, piperazin-1-yl, piperidino, pirrolidin-1-yl. The "5 to 7 membered saturated cyclic amino" can be condensed with a benzene ring.

30 Examples of "substituents" in the "5 to 7 membered saturated cyclic amino which may have substituents" include oxo, optionally halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkyl-carbonyl, optionally halogenated C₁₋₆ alkylsulfonyl, C₆₋₁₄ aryl which may have substituents, C₇₋₁₉ aralkyl which may have substituents, C₆₋₁₄ aryl-carbonyl
35 which may have substituents, 5 to 10 membered aromatic heterocyclic group which may have substituents, 5 to 8

membered monocyclic non-aromatic heterocyclic group (e.g., piperidino, piperazinyl, pyrrolidinyl, dihydropyridyl, etc.). The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 5 or more, each substituents can be the same or different.

Here, as "optionally halogenated C₁₋₆ alkyl" and "C₇₋₁₉ aralkyl which may have substituents", those exemplified as "substituents" in the above "cyclic group which may have substituents" can be used.

10 As "optionally halogenated C₁₋₆ alkyl-carbonyl" and "optionally halogenated C₁₋₆ alkylsulfonyl", those exemplified as "substituents" in the above "C₇₋₁₉ aralkyl which may have substituents" can be used.

15 Examples of the "C₆₋₁₄ aryl" in the "C₆₋₁₄ aryl which may have substituents" include phenyl, 1-naphthyl, 2-naphthyl, 2-indenyl, 2-anthryl. Phenyl is especially preferable.

As the substituents in the "C₆₋₁₄ aryl which may have substituents", those exemplified as "substituents" in the above "C₇₋₁₉ aralkyl which may have substituents" can be used.

20 The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Examples of the "C₆₋₁₄ aryl-carbonyl" in the "C₆₋₁₄ aryl-carbonyl which may have substituents" include 25 benzoyl, 1-naphthoyl, 2-naphthoyl.

As the "substituents" in the "C₆₋₁₄ aryl-carbonyl which may have substituents", those exemplified as "substituents" in the above "C₇₋₁₉ aralkyl which may have substituents" can be used. The number of substituents is, 30 for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Examples of "5 to 10 membered aromatic heterocyclic groups" in "5 to 10 membered aromatic heterocyclic groups 35 which may have substituents" include 5 to 10 membered (monocyclic or bicyclic) aromatic heterocyclic groups

containing 1 or 2 kinds of, preferably 1 to 4 hetero atoms selected from nitrogen, sulfur and oxygen atom in addition to carbon atoms. Concrete examples include 2- or 3-thienyl; 2-, 3- or 4-pyridyl; 2- or 3-furyl; 2-, 4- or 5-thiazolyl; 2-, 4- or 5-oxazolyl; 1-, 3- or 4-pyrazolyl; 2-pyrazinyl; 2-, 4- or 5-pyrimidinyl; 1-, 2- or 3-pyrrolyl; 1-, 2- or 4-imidazolyl; 3- or 4-pyridazinyl; 3-isothiazolyl; 3-isoxazolyl; 1,2,4-oxadiazol-5-yl; 1,2,4-oxadiazol-3-yl; 2-, 3-, 4-, 5- or 8-quinolyl; 1-, 3-, 4-, 5-, 6-, 7- or 8-isoquinolyl; 1-, 2-, 3-, 4-, 5-, 6- or 7-indolyl; 1-, 2-, 4- or 5-isoindolyl; 1-, 5- or 6-phthalazinyl; 2-, 3- or 5-quinoxalinyl; 2-, 3-, 4-, 5- or 6-benzofuranyl; 2-, 4-, 5- or 6-benzothiazolyl; 1-, 2-, 4-, 5- or 6-benzimidazolyl.

Examples of the "substituents" in the "5 to 10 membered aromatic heterocyclic groups which may have substituents" include halogen atom (e.g. fluorine, chlorine, bromine and iodine, etc.), C₁₋₃ alkylenedioxy (e.g. methylenedioxy, ethylenedioxy, etc.), nitro, cyano, optionally halogenated C₁₋₆ alkyl, C₆₋₁₄ aryloxy-C₁₋₆ alkyl (e.g. phenoxyethyl, etc.), C₁₋₆ alkyl-C₆₋₁₄ aryl-C₂₋₆ alkenyl (e.g. methylphenylethenyl, etc.), optionally halogenated C₃₋₆ cycloalkyl, optionally halogenated C₁₋₆ alkoxy, optionally halogenated C₁₋₆ alkylthio, C₇₋₁₉ aralkyl which may have substituents, hydroxy, C₆₋₁₄ aryloxy which may have substituents, C₇₋₁₉ aralkyloxy which may have substituents, amino, amino-C₁₋₆ alkyl (e.g. aminomethyl, aminoethyl, aminopropyl, aminobutyl, etc.), mono-C₁₋₆ alkylamino (e.g. methylamino, ethylamino, propylamino, isopropylamino, butylamino, etc.), di-C₁₋₆ alkylamino (e.g. dimethylamino, diethylamino, dipropylamino, dibutylamino, ethylmethylamino, etc.), mono-C₁₋₆ alkylamino-C₁₋₆ alkyl (e.g. methylaminomethyl, ethylaminomethyl, propylaminomethyl, isopropylaminoethyl, butylaminoethyl, etc.), di-C₁₋₆ alkylamino-C₁₋₆ alkyl (e.g. dimethylaminomethyl, diethylaminomethyl,

dipropylaminomethyl, diisopropylaminoethyl, dibutylaminoethyl, etc.), 5 to 7 membered saturated cyclic amino, acyl, acylamino, acyloxy. The number of substituents is, for instance, 1 to 5, preferably 1 to 3.

5 When the number of substituents is 2 or more, each substituents can be the same or different.

Here, as "optionally halogenated C₁₋₆ alkyl", "optionally halogenated C₃₋₆ cycloalkyl", "optionally halogenated C₁₋₆ alkoxy", "optionally halogenated C₁₋₆ alkylthio", "C₇₋₁₁, aralkyl which may have substituents", "C₆₋₁₄ aryloxy which may have substituents", "C₇₋₁₉, aralkyloxy which may have substituents", those exemplified as the "substituent" in the above "cyclic group which may have substituents" can be used respectively.

15 As a "5 to 7 membered saturated cyclic amino", those exemplified as "5 to 7 membered saturated cyclic amino" regarding "5 to 7 membered saturated cyclic amino which may have substituents" which is a "substituent" in the above "5 to 7 membered saturated cyclic amino which may have 20 substituents" can be used.

Examples of the above "acyl" include acyl of the formulae : -CO-R³, -CO-OR³, -CO-NR³R⁴, -CS-NR³R⁴, -SO₂-R^{3a}, -SO-R^{3a}, -PO(-OR³)-OR⁴ or -PO₂-R^{3a} wherein R³ is (i) hydrogen atom, (ii) a hydrocarbon group which may have 25 substituents, or (iii) a heterocyclic group which may have substituents; R^{3a} is (i) a hydrocarbon group which may have substituents, or (ii) a heterocyclic group which may have substituents; R⁴ is hydrogen atom or C₁₋₆ alkyl; R³ and R^{3a}, together with the adjacent nitrogen atom, can form a 30 nitrogen-containing hetero ring which may have substituents.

Examples of the "hydrocarbon group" in "hydrocarbon group which may have substituents" for R³ or R⁴ include straight-chain or cyclic hydrocarbon groups (e.g. alkyl, 35 alkenyl, alkynyl, cycloalkyl, aryl, aralkyl, etc.). Among these, C₁₋₁₉ straight chain or cyclic hydrocarbon groups as

shown below are preferable.

a) C₁₋₆ alkyl (e.g. methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, hexyl, etc.);

5 b) C₂₋₆ alkenyl (e.g., vinyl, allyl, isopropenyl, 2-butenyl, etc.);

c) C₂₋₆ alkynyl (e.g. ethynyl, propargyl, 2-butynyl, etc.);

d) C₃₋₆ cycloalkyl (e.g. cyclopropyl, cyclobutyl, 10 cyclopentyl, cyclohexyl, etc.); the C₃₋₆ cycloalkyl can be condensed with one benzene ring;

e) C₆₋₁₄ aryl (e.g. phenyl, 1-naphthyl, 2-naphthyl, 2-indenyl, 2-anthryl, etc.), preferably phenyl;

f) C₇₋₁₉ aralkyl (e.g. benzyl, phenethyl, 15 diphenylmethyl, triphenylmethyl, 1-naphthylmethyl, 2-naphthylmethyl, 2,3-diphenylethyl, 3-phenylpropyl, 4-phenylbutyl, 5-phenylpentyl, etc.), preferably benzyl.

The "hydrocarbon groups" are preferably C₁₋₆ alkyl, C₆₋₁₄ aryl, C₇₋₁₉ aralkyl, etc.

20 Examples of the "substituent" in "hydrocarbon groups which may have substituents" include halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), C₁₋₃ alkylenedioxy (e.g. methylenedioxy, ethylenedioxy, etc.), 25 nitro, cyano, optionally halogenated C₁₋₆ alkoxy, optionally halogenated C₁₋₆ alkylthio, hydroxy, amino, mono-C₁₋₆ alkylamino (e.g. methylamino, ethylamino, propylamino, isopropylamino, butylamino, etc.), di-C₁₋₆ alkylamino (e.g. dimethylamino, diethylamino, 30 dipropylamino, dibutylamino, ethylmethyleamino, etc.), formyl, carboxy, carbamoyl, thiocarbamoyl, optionally halogenated C₁₋₆ alkyl-carbonyl, C₁₋₆ alkoxy-carbonyl (e.g., methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, tert-butoxycarbonyl, etc.), 5 to 10 membered aromatic 35 heterocyclic groups which may have substituents, C₆₋₁₄ aryl-carbonyl which may have substituents, C₆₋₁₄

aryloxy-carbonyl which may have substituents, C₇₋₁₉, aralkyloxy-carbonyl which may have substituents, 5 to 6 membered hetero ring-carbonyl which may have substituents, mono-C₁₋₆ alkyl-carbamoyl (e.g. methylcarbamoyl, 5 ethylcarbamoyl, etc.), di-C₁₋₆ alkyl-carbamoyl (e.g. dimethylcarbamoyl, diethylcarbamoyl, ethylmethylcarbamoyl, etc.), C₆₋₁₄ aryl-carbamoyl which may have substituents, 5 to 6 membered hetero ring-carbamoyl which may have substituents, optionally halogenated C₁₋₆, 10 alkylsulfonyl, C₆₋₁₄ arylsulfonyl which may have substituents, formylamino, C₁₋₆ alkyl-carbonyloxy (e.g. acetoxy, propanoyloxy, etc.), C₆₋₁₄ aryl-carbonyloxy which may have substituents, C₁₋₆ alkoxy-carbonyloxy (e.g. methoxycarbonyloxy, ethoxycarbonyloxy, 15 propoxycarbonyloxy, butoxycarbonyloxy, etc.), mono-C₁₋₆ alkyl-carbamoyloxy (e.g. methylcarbamoyloxy, ethylcarbamoyloxy, etc.), di-C₁₋₆ alkyl-carbamoyloxy (e.g. dimethylcarbamoyloxy, diethylcarbamoyloxy, etc.), C₆₋₁₄ aryl-carbamoyloxy which may have substituents, 20 nicotinoyloxy. The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Here, as "optionally halogenated C₁₋₆ alkoxy", 25 "optionally halogenated C₁₋₆ alkylthio" and "C₆₋₁₄ aryl-carbamoyl which may have substituents", those exemplified as a "substituent" in the above "cyclic group which may have substituents" can be used.

As "optionally halogenated C₁₋₆ alkyl-carbonyl" and 30 "optionally halogenated C₁₋₆ alkylsulfonyl", those exemplified as a "substituent" in the above "C₇₋₁₉, aralkyl which may have substituents" can be used.

As the above "5 to 10 membered aromatic heterocyclic groups which may have substituents" and "C₆₋₁₄ aryl-carbonyl 35 which may have substituents", those exemplified as "substituent" in the above "5 to 7 membered saturated cyclic

amino which may have substituents" can be used.

Examples of "C₆₋₁₄ aryloxy-carbonyl" in "C₆₋₁₄ aryloxy-carbonyl which may have substituents" include phenyloxycarbonyl, 1-naphthyloxycarbonyl, 2-
5 naphthyloxycarbonyl.

Examples of "C₇₋₁₉ aralkyloxy-carbonyl" in "C₇₋₁₉ aralkyloxy-carbonyl which may have substituents" include benzyloxycarbonyl, phenethyloxycarbonyl,
diphenylmethyloxycarbonyl, triphenylmethyloxycarbonyl,
10 1-naphthylmethyloxycarbonyl, 2-
naphthylmethyloxycarbonyl, 2,2-
diphenylethyloxycarbonyl, 3-phenylpropyloxycarbonyl, 4-
phenylbutyloxycarbonyl, 5-phenylpentyloxycarbonyl.

Examples of "5 to 6 membered hetero ring-carbonyl" in
15 the above "5 to 6 membered hetero ring-carbonyl which may
have substituents" include nicotinoyl, isonicotinoyl,
2-thenoyl, 3-thenoyl, 2-furoyl, 3-furoyl,
morpholinocarbonyl, pepiridinocarbonyl, pyrrolidin-1-
ylcarbonyl.

20 Examples of the "5 to 6 membered hetero ring-
carbamoyl" in the above "5 to 6 membered hetero ring-
carbamoyl which may have substituents" include
morpholinocarbamoyl, pepiridinocarbamoyl, 2-
pyridylcarbamoyl, 3-pyridylcarbamoyl, 4-
25 pyridylcarbamoyl, 2-thienylcarbamoyl, 3-
thienylcarbamoyl.

Examples of "C₆₋₁₄ arylsulfonyl" in the above "C₆₋₁₄ arylsulfonyl which may have substituents" include phenylsulfonyl, 1-naphthylsulfonyl, 2-naphthylsulfonyl.

30 Examples of "C₆₋₁₄ aryl-carbonyloxy" in the above "C₆₋₁₄ aryl-carbonyloxy which may have substituents" include benzoxyloxy, 1-naphthoxyloxy, 2-naphthoxyloxy.

Examples of "C₆₋₁₄ aryl-carbamoyloxy" in the above "C₆₋₁₄ aryl-carbamoyloxy which may have substituents" include phenylcarbamoyloxy, naphthylcarbamoyloxy.

As the "substituents" in the above "C₆₋₁₄ aryloxy-

carbonyl which may have substituents", "C₇₋₁₉, aralkyloxy-carbonyl which may have substituents", "5 to 6 membered hetero ring-carbonyl which may have substituents", "5 to 6 membered hetero ring-carbamoyl which 5 may have substituents", "C₆₋₁₄ arylsulfonyl which may have substituents", "C₆₋₁₄ aryl-carbonyloxy which may have substituents" and "C₆₋₁₄ aryl-carbamoyloxy which may have substituents", those exemplified as "substituents" in the above "C₇₋₁₉, aralkyl which may have substituents" can be 10 mentioned. The number of the substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

15 Examples of "heterocyclic groups" in the "heterocyclic groups which may have substituents" for R³ or R^{3a} include a 5 to 14 membered (monocyclic, bicyclic or tricyclic) hetero ring containing 1 or 2 kinds of, 1 to 4 hetero atoms selected from nitrogen, sulfur and oxygen atom 20 in addition to carbon atoms. Preferably, univalent groups formed by removing an optional one hydrogen atom from (i) an aromatic hetero ring, (ii) a 5 to 10 membered non-aromatic hetero ring, or (iii) a 7 to 10 membered hetero-bridge ring, can be mentioned.

25 Here, examples of the "aromatic hetero ring" include a 5 to 14 membered, preferably 5 to 10 membered, aromatic hetero ring containing one or more hetero atom (e.g. 1 to 4) selected from nitrogen, sulfur and oxygen atom in addition to carbon atoms.

30 Concrete examples include aromatic hetero rings such as thiophene, furan, pyrrole, imidazole, pyrazole, thiazole, isothiazole, oxazole, isoxazole, pyridine, pyrazine, pyrimidine, pyridazine, 1,2,4-oxadiazole, 1,3,4-oxadiazole, 1,2,4-thiadiazole, 1,3,4-thiadiazole, 35 furazan, benzothiophene, benzofuran, benzimidazole, benzoxazole, benzothiazole, benzisothiazole,

naphtho[2,3-b]thiophene, phenoxathiin, indole,
isoindole, 1H-indazole, purine, 4H-quinolidine,
isoquinoline, quinoline, phthalazine, naphthyridine,
quinoxaline, quinazoline, cinnoline, carbazole, β -
5 carboline, phenanthridine, acridine,
phenazinephenothiadine, phenoxazine, phthalimide, etc.;
or a ring formed by condensing these rings (preferably
monocyclic rings) with one to multiple (preferably 1 or 2)
aromatic rings (e.g. benzene ring, etc.).

10 Examples of "5 to 10 membered non-aromatic hetero
rings" include 2- or 3-pyrroline, pyrrolidine, 2- or 3-
imidazoline, 2-oxazoline, oxazolidine, 2- or 3-pyrazoline,
pyrazolidine, 2-thiazoline, piperidine, piperazine,
hexamethylenimine, morpholine, thiomorpholine.

15 Examples of "7 to 10 membered hetero-bridge rings"
include quinuclidine, 7-azabicyclo[2.2.1]heptane.

The "hetero cyclic groups" are preferably 5 to 10
membered (monocyclic or bicyclic) heterocyclic groups
containing 1 or 2 kinds of, preferably 1 to 4, hetero atoms
20 selected from nitrogen, sulfur and oxygen atom in addition
to carbon atoms. Concretely examples include aromatic
heterocyclic groups such as 2- or 3-thienyl; 2-, 3- or
4-pyridyl; 2- or 3-furyl; 2-, 4- or 5-thiazolyl; 2-, 4- or
5-oxazolyl; 1-, 3- or 4-pyrazolyl; 2-pyrazinyl; 2-, 4- or
25 5-pyrimidinyl; 1-, 2- or 3-pyrrolyl; 1-, 2- or 4-
imidazolyl; 3- or 4-pyridazinyl; 3-isothiazolyl; 3-
isoxazolyl; 1,2,4-oxadiazol-5-yl; 1,2,4-oxadiazol-3-yl;
2-, 3-, 4-, 5- or 8-quinolyl; 1-, 3-, 4-, 5-, 6-, 7- or
8-isoquinolyl; 1-, 2-, 3-, 4-, 5-, 6- or 7-indolyl; 1-, 2-,
30 4- or 5-isoindolyl; 1-, 5- or 6-phthalazinyl; 2-, 3- or
5-quinoxalinyl; 2-, 3-, 4-, 5- or 6-benzofuranyl; 2-, 3-,
4-, 5- or 6-benzothienyl; 2-, 4-, 5- or 6-benzothiazolyl;
1-, 2-, 4-, 5- or 6-benzimidazolyl; and non-aromatic
heterocyclic groups such as 1-, 2- or 3-pyrrolidinyl; 1-,
35 2-, 4- or 5-imidazolidinyl; 2- or 4-imidazolinyl; 2-, 3-
or 4-pyrazolidinyl; piperidino; 2-, 3- or 4-piperidyl; 1-

or 2-piperazinyl; morpholino.

As the "substituents" in the "heterocyclic groups which may have substituents", those exemplified as "substituents" in the above "5 to 10 membered aromatic heterocyclic groups which may have substituents" can be used. The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Examples of "C₁₋₆ alkyl" for R⁴ include methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, hexyl.

Examples of "nitrogen-containing hetero ring" in the "nitrogen-containing hetero ring which may have substituents" formed by R³ and R⁴ together with the adjacent nitrogen atoms, include a 5 to 7 membered nitrogen-containing hetero ring which contains at least one nitrogen atom in addition to carbon atoms and may contain 1 to 3 hetero atoms selected from nitrogen, sulfur and oxygen atom. The "nitrogen-containing hetero rings" are preferably piperidine, morpholine, thiomorpholine, piperazine, pyrrolidine, etc.

As the "substituents" in the "nitrogen-containing hetero ring which may have substituents", those exemplified as "substituents" in the above "5 to 10 membered aromatic heterocyclic groups which may have substituents" can be used. The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

The "acyl" is preferably formyl, carboxy, carbamoyl, optionally halogenated C₁₋₆ alkyl-carbonyl (e.g. acetyl, etc.), C₁₋₆ alkoxy-carbonyl (e.g. methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, tert-butoxycarbonyl, etc.), C₆₋₁₄ aryl-carbonyl which may have substituents (e.g. benzoyl, 1-naphthoyl, 2-naphthoyl, etc.), C₆₋₁₄ aryloxy-carbonyl which may have substituents (e.g. phenyloxycarbonyl, 1-naphthyloxycarbonyl, 2-

naphthyoxy carbonyl; etc.), C_{7-19} aralkyloxy-carbonyl which may have substituents (e.g. benzyloxy carbonyl, phenethyloxy carbonyl, etc.), a 5 to 6 membered hetero ring-carbonyl which may have substituents (e.g. 5 nicotinoyl, etc.), mono- C_{1-6} alkyl-carbamoyl (e.g. methylcarbamoyl, ethylcarbamoyl, etc.), di- C_{1-6} alkyl-carbamoyl (e.g. dimethylcarbamoyl, diethylcarbamoyl, ethylmethylcarbamoyl, etc.), C_{6-14} aryl-carbamoyl which may have substituents (e.g. phenylcarbamoyl, 4-methoxyphenylcarbamoyl, 3,4-dimethoxyphenylcarbamoyl, etc.), aromatic hetero ring-carbamoyl which may have substituents (e.g. 2-pyridinylcarbamoyl, 2-quinolinylcarbamoyl etc.), optionally halogenated C_{1-6} alkylsulfonyl (e.g. methylsulfonyl, etc.), C_{6-14} arylsulfonyl which may have substituents (e.g. phenylsulfonyl etc.), etc.

Here, as "optionally halogenated C_{1-6} alkyl-carbonyl" and "optionally halogenated C_{7-19} aralkylsulfonyl", those exemplified as "substituents" in the above " C_{7-19} aralkyl which may have substituents" can be used.

As " C_{6-14} aryl-carbonyl which may have substituents", "substituents" in the above "5 to 7 membered saturated cyclic amino which may have substituents" can be used.

As " C_{6-14} aryloxy-carbonyl which may have substituents", " C_{7-19} aralkyloxy-carbonyl which may have substituents", "5 to 6 membered hetero ring-carbonyl which may have substituents", "aromatic hetero ring-carbamoyl which may have substituents" and " C_{6-14} arylsulfonyl which may have substituents", those exemplified as "substituents" in the above "hydrocarbon groups which may have substituents" can be used.

As " C_{6-14} aryl-carbamoyl which may have substituents", those exemplified as "substituents" in the above "cyclic group which may have substituents" can be used.

is substituted by 1 or 2 of the above "acyl". Preferably, acylamino of the formulae : -NR⁵-COR⁶, -NR⁵-COOR^{6a}, -NR⁵-SO₂R^{6a}, -NR⁵-CONR^{6a}R^{6b}, -PO(-OR⁵)-OR⁶, or -PO₂-R⁶ wherein R⁵ is hydrogen atom or C₁₋₆ alkyl; R⁶ has the same meaning as 5 the above R³; R^{6a} has the same meaning as the above R^{3a}; and R^{6b} has the same meaning as R⁴], can be mentioned.

As "C₁₋₆ alkyl" for R⁵, the same one as in "C₁₋₆ alkyl" for the above R⁴ can be mentioned.

The "acylamino" is preferably formylamino, optionally 10 halogenated C₁₋₆ alkyl-carboxamide (e.g. methylcarboxamide, trifluoromethylcarboxamide, isopropylcarboxamide, etc.), C₆₋₁₄ aryl-carboxamide which may have substituents (e.g. phenylcarboxamide, 2-methoxyphenylcarboxamide, 4-methoxyphenylcarboxamide, etc.), N-(C₆₋₁₄ aryl-carbonyl which may have 15 substituents)-N-C₁₋₆ alkylamino (e.g. N-4-methoxybenzoyl-N-methylamino, etc.), C₇₋₁₉ aralkyl-carboxamide which may have substituents (e.g. benzylcarboxamide, etc.), aromatic hetero ring-20 carboxamide which may have substituents (e.g. benzothiophen-2-ylcarboxamide, etc.), optionally halogenated C₁₋₆ alkoxy-carboxamide (e.g. methoxycarboxamide, ethoxycarboxamide, propoxycarboxamide, butoxycarboxamide, etc.), C₆₋₁₄ 25 arylamino-carbonylamino which may have substituents (e.g. phenylaminocarbonylamino, etc.), optionally halogenated C₁₋₆ alkylsulfonylamino (e.g. methylsulfonylamino, trifluoromethylsulfonylamino, ethylsulfonylamino, etc.), C₆₋₁₄ arylsulfonylamino which may have substituents (e.g. 30 4-methoxyphenylsulfonylamino, etc.).

Here, as "substituents" in "C₆₋₁₄ aryl-carboxamide which may have substituents", "N-(C₆₋₁₄ aryl-carbonyl which may have substituents)-N-C₁₋₆ alkylamino", "C₇₋₁₉ aralkyl-carboxamide which may have substituents", 35 "aromatic hetero ring-carboxamide which may have substituents", "C₆₋₁₄ arylamino-carbonylamino which may

have substituents" and "C₆₋₁₄ arylsulfonylamino which may have substituents", those exemplified as "substituents" in the above "C₇₋₁₉ aralkyl which may have substituents" can be mentioned. The number of substituents is, for instance,
5 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Examples of the above "acyloxy" include oxy substituted by one of the above "acyl". Preferably,
10 acyloxy of the formulae : -O-COR⁷, -O-COOR⁷, -O-CONHR⁷, -PO(OH)-OR⁷ or -PO₂-R⁷ wherein R⁷ has the same meaning as the above R³, can be mentioned.

The "acyloxy" is preferably optionally halogenated C₁₋₆ alkyl-carbonyloxy (e.g. acetoxy, propanoyloxy, etc.),
15 C₆₋₁₄ aryl-carbonyloxy which may have substituents (e.g. benzyloxy, 4-methoxybenzyloxy, etc.), optionally halogenated C₁₋₆ alkoxy-carbonyloxy (e.g. methoxycarbonyloxy, trifluoromethoxycarbonyloxy, ethoxycarbonyloxy, propoxycarbonyloxy,
20 butoxycarbonyloxy, etc.), mono-C₁₋₆ alkyl-carbamoyloxy (e.g. methylcarbamoyloxy, ethylcarbamoyloxy, etc.), di-C₁₋₆ alkyl-carbamoyloxy (e.g. dimethylcarbamoyloxy, diethylcarbamoyloxy, etc.), C₆₋₁₄ aryl-carbamoyloxy which may have substituents (e.g. phenylcarbamoyloxy,
25 naphthylcarbamoyloxy, etc.), nicotinyloxy, etc.

As "substituents" in "C₆₋₁₄ aryl-carbonyloxy which may have substituents" and "C₆₋₁₄ aryl-carbamoyloxy which may have substituents", those exemplified as "substituents" in the above "C₇₋₁₉ aralkyl which may have substituents" can be mentioned. The number of substituents is, for instance,
30 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

Examples of the "5 to 7 membered non-aromatic
35 heterocyclic groups which may have substituents", which is "substituents" in "cyclic group which may have

substituents" for Ar¹, include 4,5-dihydro-1,3-oxazol-2-yl, 4,5-dihydro-1,3-thiazol-2-yl, 4,5-dihydro-1H-2-imidazolyl. As "substituents" in the "5 to 7 membered non-aromatic heterocyclic groups which may have substituents", those exemplified as "substituents" in the above "5 to 7 membered saturated cyclic amino which may have substituents" can be used.

5 As "acyl", "acyloxy" and "acylamino", which are "substituents" in the "cyclic group which may have substituents" for Ar¹, those exemplified as "substituents" in the above "5 to 10 membered aromatic heterocyclic groups which may have substituents" can be used.

10 Regarding "aromatic hetero ring-C₁₋₆ alkoxy" which is "substituents" in the "cyclic group which may have substituents" for Ar¹, as "aromatic hetero ring", those exemplified as the above R³ can be used. Examples of "C₁₋₆ alkoxy" include methoxy, ethoxy, propoxy, butoxy, pentyloxy.

15 20 "Substituents" in the "cyclic group which may have substituents" for Ar¹ are preferably halogen atom (preferably fluorine, chlorine and bromine, etc.); nitro; C₁₋₃ alkylenedioxy (preferably methylenedioxy, etc.); optionally halogenated C₁₋₆ alkyl (preferably, methyl, ethyl, propyl, trifluoromethyl, etc.); hydroxy-C₁₋₆ alkyl (preferably hydroxymethyl, etc.); optionally halogenated C₃₋₆ cycloalkyl (preferably cyclohexyl, etc.); optionally halogenated C₁₋₆ alkoxy (preferably methoxy, ethoxy, etc.); optionally halogenated C₁₋₆ alkylthio (preferably 25 methylthio, etc.); hydroxy; C₇₋₁₁ aralkyloxy which may have substituents (preferably, 1 to 3 substituents selected from halogen atom, optionally halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkoxy, optionally halogenated C₁₋₆ alkylthio, etc.) (preferably benzyloxy, 4-30 methoxybenzyloxy, 3-methoxybenzyloxy, 4-fluorobenzyloxy, 4-methylthiobenzyloxy, 4-ethylbenzyloxy, etc.); C₆₋₁₄

aryloxy which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably phenoxy, 4-methoxyphenoxy, etc.); amino; mono-C₁₋₆ alkylamino (preferably methylamino, etc.); di-C₁₋₆

5 alkylamino (preferably dimethylamino, etc.); 5 to 7 membered saturated cyclic amino which may have substituents (preferably 1 to 3 oxo) and may be condensed with a benzene ring (preferably 1,3-dioxo-1,3-dihydro-2H-isoindol-2-yl, etc.); 5 to 7 membered non-aromatic heterocyclic groups

10 which may have substituents (preferably 4,5-dihydro-1,3-oxazol-2-yl, etc.); formyl; carboxy; C₆₋₁₄ aryl-carbonyl which may have substituents (preferably benzoyl, etc.); C₆₋₁₄ aryl-carbamoyl which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.)

15 (preferably, phenylcarbamoyl, 4-methoxyphenylcarbamoyl, 3,4-dimethoxyphenylcarbamoyl, etc.); aromatic hetero ring-carbamoyl which may have substituents (preferably 2-pyridinylcarbamoyl, 2-quinolinylcarbamoyl, etc.); C₁₋₆ alkoxy-carbonyl (preferably methoxycarbonyl,

20 ethoxycarbonyl, etc.); optionally halogenated C₁₋₆ alkyl-carboxamide (preferably methylcarboxamide, trifluoromethylcarboxamide, isopropylcarboxamide, etc.); C₆₋₁₄ aryl-carboxamide which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.)

25 (preferably phenylcarboxamide, 2-methoxyphenylcarboxamide, 4-methoxyphenylcarboxamide, etc.); C₇₋₁₉ aralkyl-carboxamide which may have substituents (preferably benzylcarboxamide, etc.); aromatic hetero ring-carboxamide which may have substituents (preferably

30 benzothiophen-2-ylcarboxamide, etc.); N-(C₆₋₁₄ aryl-carbonyl which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.))-N-C₁₋₆ alkylamino (preferably N-4-methoxybenzoyl-N-methylamino, etc.); C₆₋₁₄ arylamino-carbonylamino which may have substituents

35 (preferably phenylaminocarbonylamino, etc.); C₆₋₁₄ arylsulfonylamino which may have substituents (preferably,

1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably 4-methoxyphenylsulfonylamino, etc.); C₆₋₁₄ aryl carbonyloxy which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably 5 4-methoxybenzoyloxy, etc.); oxo; carboxy-C₁₋₆ alkyl (preferably carboxyethyl, etc.); C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl (preferably methoxycarbonylmethyl, etc.); C₇₋₁₉ aralkyl which may have substituents (preferably 1 to 3 halogen atom) (preferably benzyl, 2,4-dichlorobenzyl, 10 etc.); aromatic hetero ring-C₁₋₆ alkoxy (preferably 2-qunolylmethoxy, etc.); cyano, etc.

When "cyclic group" in "cyclic group which may have substituents" for Ar¹ is a non-aromatic cyclic hydrocarbon group or a non-aromatic heterocyclic group, C₆₋₁₄ aryl which 15 may have substituents (preferably, 1 to 3 substituents selected from halogen atom, C₁₋₃ alkylenedioxy, optionally halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkoxy, etc.) (preferably phenyl, 4-fluorophenyl, 1,3-benzodioxol-5-yl, 4-chlorophenyl, 4-methylphenyl, 4-methoxyphenyl), hydroxy, C₇₋₁₉, aralkyloxy-carbonyl (preferably benzyloxycarbonyl), C₇₋₁₉, aralkyl (preferably benzyl), etc., can be used as a preferable substituent.

Ar¹ is preferably phenyl, biphenylyl (preferably 25 4-biphenylyl, 2-biphenylyl), phenyl-pyridyl (preferably 6-phenyl-3-pyridyl, 5-phenyl-2-pyridyl), phenyl-furyl (preferably 5-phenyl-2-furyl), phenyl-isoxazolyl (preferably 3-phenyl-isoxazol-5-yl), diphenyl-oxazolyl (preferably 2,4-diphenyl-1,3-oxazol-5-yl), pyridyl-30 phenyl (preferably 4-(4-pyridyl)phenyl, 4-(3-pyridyl)phenyl), phenyl-pyrimidinyl (preferably 2-phenyl-5-pyrimidinyl), benzofuranyl-phenyl (preferably 4-(2-benzofuranyl)phenyl), furyl-phenyl (preferably 4-(2-furyl)phenyl), terphenyl (preferably 4,4'-terphenyl), 35 thienyl-phenyl (preferably 4-(2-thienyl)phenyl), indolyl (preferably 2-indolyl, 3-indolyl), naphthyl-oxadiazolyl

(preferably 3-(2-naphthyl)-1,2,4-oxadiazol-5-yl), benzofuranyl-oxadiazole (preferably 3-(2-benzofuranyl)-1,2,4-oxadiazol-5-yl), benzothienyl (preferably 2-benzothienyl), benzofuranyl (preferably 2-benzofuranyl),
5 fluorenyl (preferably 2-fluorenyl), pyridyl-pyrrolyl (preferably 3-(4-pyridyl)pyrrolyl), thioxanthenyl; each of which may have 1 to 3 (preferably 1 or 2) substituents selected from the group consisting of halogen atom (preferably fluorine, chlorine, bromine, etc.); nitro; C₁₋₃ alkylenedioxy (preferably methylenedioxy, etc.); optionally halogenated C₁₋₆ alkyl (preferably methyl, ethyl, propyl, trifluoromethyl, etc.); hydroxy-C₁₋₆ alkyl (preferably hydroxymethyl, etc.); optionally halogenated C₃₋₆ cycloalkyl (preferably cyclohexyl, etc.); optionally 10 halogenated C₁₋₆ alkoxy (preferably methoxy, ethoxy, etc.); optionally halogenated C₁₋₆ alkythio (preferably methylthio, etc.); hydroxy; C₇₋₁₉ aralkyloxy which may have substituents (preferably, 1 to 3 substituents selected from halogen atom, optionally halogenated C₁₋₆ alkyl, optionally 15 halogenated C₁₋₆ alkoxy, optionally halogenated C₁₋₆ alkylthio, etc.) (preferably benzyloxy, 4-methoxybenzyloxy, 3-methoxybenzyloxy, 4-fluorobenzyloxy, 4-methylthiobenzyloxy, 4-ethylbenzyloxy, etc.); C₆₋₁₄ aryloxy which may have substituents (preferably, 1 to 3 20 substituents selected from halogen atom, optionally halogenated C₁₋₆ alkoxy, etc.) (preferably phenyloxy, 4-methoxyphenyloxy, etc.); amino; mono-C₁₋₆ alkylamino (preferably methylamino, etc.); di-C₁₋₆ alkylamino (preferably dimethylamino, etc.); 5 to 7 membered saturated cyclic amino which may have substituents 25 (preferably 1 to 3 oxo) and may be condensed with a benzene ring (preferably 1,3-dioxo-1,3-dihydro-2H-isoindol-2-yl, etc.); 5 to 7 membered non-aromatic heterocyclic groups which may have substituents (preferably 4,5-dihydro-1,3-oxazol-2-yl, etc.); formyl; carboxy; C₆₋₁₄ aryl-30 carbonyl which may have substituents (preferably benzoyl, etc.); C₆₋₁₄ aryl-carbamoyl which may have substituents 35

(preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably phenylcarbamoyl, 4-methoxyphenylcarbamoyl, 3,4-dimethoxyphenylcarbamoyl, etc.); aromatic hetero ring-carbamoyl which may have substituents (e.g. 2-5 pridinylcarbamoyl, 2-quinolinylcarbamoyl, etc.); C₁₋₆ alkoxy-carbonyl (preferably methoxycarbonyl, ethoxycarbonyl, etc.); optionally halogenated C₁₋₆ alkyl-carboxamide (preferably, methylcarboxamide, trifluoromethylcarboxamide, isopropylcarboxamide, etc.); 10 C₆₋₁₄ aryl-carboxamide which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably phenylcarboxamide, 2-methoxyphenylcarboxamide, 4-methoxyphenylcarboxamide, etc.); C₇₋₁₉, aralkyl-carboxamide which may have substituents 15 (preferably benzylcarboxamide, etc.); aromatic hetero ring-carboxamide which may have substituents (preferably benzothiophen-2-ylcarboxamide, etc.); N-(C₆₋₁₄ aryl-carbonyl which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.))-N-C₁₋₆ alkylamino 20 (preferably N-4-methoxybenzoyl-N-methylamino, etc.); C₆₋₁₄ arylamino-carbonylamino which may have substituents (preferably phenylaminocarbonylamino, etc.); C₆₋₁₄ arylsulfonylamino which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably 25 4-methoxyphenylsulfonylamino, etc.); C₆₋₁₄ aryl-carbonyloxy which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably 4-methoxybenzoyloxy, etc.); oxo; carboxy-C₁₋₆ alkyl (preferably carboxyethyl, etc.); C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl (preferably methoxycarbonylmethyl, etc.); C₇₋₁₉, aralkyl which may have substituents (preferably 1 to 3 halogen atom) (preferably benzyl, 2,4-dichlorobenzyl, etc.); aromatic hetero ring-C₁₋₆ alkoxy (preferably 2-qunolylmethoxy, etc.); and cyano.

30 Further, preferable examples of Ar¹ include piperidinyl (preferably piperidino), piperazinyl,

pyrrolidinyl, dihydropyridyl, tetrahydropyridyl; each of which may have 1 or 2 substituents selected from the group consisting of oxo, C₆₋₁₄ aryl which may have substituents (preferably, 1 to 3 substituents selected from halogen atom, C₁₋₃ alkyleneoxy, optionally halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkoxy, etc.) (preferably phenyl, 4-fluorophenyl, 1,3-benzodioxol-5-yl, 4-chlorophenyl, 4-methylphenyl, 4-methoxyphenyl), hydroxy, C₇₋₁₉, aralkyloxy-carbonyl (preferably benzyloxycarbonyl) and C₇₋₁₉, aralkyl (preferably benzyl).

Ar¹ is more preferably, phenyl, biphenylyl (preferably 4-biphenylyl) or phenyl-pyridyl (preferably 6-phenyl-3-pyridyl, 5-phenyl-2-pyridyl); each of which may have 1 or 2 substituents selected from the group consisting of halogen atom (preferably fluorine, chlorine, bromine, etc.); optionally halogenated C₁₋₆ alkyl (preferably methyl, ethyl, propyl, trifluoromethyl, etc.); optionally halogenated C₁₋₆ alkoxy (preferably methoxy, ethoxy, etc.); C₇₋₁₉, aralkyloxy which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkoxy, optionally halogenated C₁₋₆ alkylthio, etc.) (preferably benzyloxy, 4-methoxybenzyloxy, etc.); C₆₋₁₄ aryloxy which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably phenoxy, etc.); C₆₋₁₄ aryl-carbonyl which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably benzoyl, etc.); C₆₋₁₄ aryl-carbamoyl which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably phenylcarbamoyl, 4-methoxyphenylcarbamoyl, 3,4-dimethoxyphenylcarbamoyl, etc.); aromatic hetero ring-carbamoyl which may have substituents (e.g. 2-pyridinylcarbamoyl, 2-quinolinylcarbamoyl, etc.); C₆₋₁₄ aryl-carboxamide which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably

phenylcarboxamide, 2-methoxyphenylcarboxamide, 4-methoxyphenylcarboxamide, etc.); C₇₋₁₉ aralkyl-carboxamide which may have substituents (preferably benzylcarboxamide, etc.); aromatic hetero ring-carboxamide (preferably 5 benzothiophen-2-ylcarboxamide, etc.); N-(C₆₋₁₄ aryl-carbonyl which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.))-N-C₁₋₆ alkylamino (preferably N-4-methoxybenzoyl-N-methylamino, etc.); C₆₋₁₄ arylamino-carbonylamino which may have substituents 10 (preferably phenylaminocarbonylamino, etc.); C₆₋₁₄ arylsulfonylamino which may have substituents (preferably, 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably 4-methoxyphenylsulfonylamino, etc.); and C₆₋₁₄ arylcarbonyloxy which may have substituents (preferably, 15 1 to 3 optionally halogenated C₁₋₆ alkoxy, etc.) (preferably 4-methoxybenzoyloxy, etc.).

Further, preferable examples of Ar¹ include 20 piperidino, piperazinyl or pyrrolidinyl; each of which may have 1 or 2 substituents selected from the group consisting of oxo and C₆₋₁₄ aryl (preferably phenyl) which may have substituents [preferably halogen atom (preferably fluorine, chlorine, bromine, etc.), optionally halogenated C₁₋₆ alkyl (preferably methyl, ethyl, propyl, trifluoromethyl, etc.) or optionally halogenated C₁₋₆ 25 alkoxy (preferably methoxy, ethoxy, etc.)].

The "spacer having a main chain of 1 to 6 atoms" means a space in which 1 to 6 atoms are linked. Here, the "number of atoms in the main chain" is counted so that the number 30 of atoms in the main chain is minimum. For instance, the number of atoms of 1,2-cyclopentylene is counted as 2, and the number of atoms of 1,3-cyclopentylene is counted as 3.

Examples of the "spacer having a main chain of 1 to 6 atoms" include a bivalent group consisting of 1 to 3 35 species selected from -O-, -S-, -CO-, -SO-, -SO₂-, -NR⁸- (R⁸ is hydrogen atom, optionally halogenated C₁₋₆ alkyl,

optionally halogenated C₁₋₆ alkyl-carbonyl, optionally halogenated C₁₋₆ alkylsulfonyl), bivalent C₁₋₆ non-cyclic hydrocarbon groups which may have substituents, and bivalent C₅₋₈ monocyclic non-aromatic hydrocarbon groups.

5 Here, as "optionally halogenated C₁₋₆ alkyl", those exemplified as "substituents" in the above "cyclic group which may have substituents" can be used.

10 As "optionally halogenated C₁₋₆ alkyl-carbonyl" and "optionally halogenated C₁₋₆ alkylsulfonyl", those exemplified as "substituents" in the above "C₇₋₁₉, aralkyl which may have substituents" can be used.

15 Examples of "bivalent C₁₋₆ non-cyclic hydrocarbon groups" in the "bivalent C₁₋₆ non-cyclic hydrocarbon groups which may have substituents" include

(1) C₁₋₆ alkylene (e.g. -CH₂-, -(CH₂)₂-, -(CH₂)₃-, -(CH₂)₄-, -(CH₂)₅-, -(CH₂)₆-, -CH(CH₃)₂-, -C(CH₃)₂-, -(CH(CH₃))₂-, -(CH₂)₂C(CH₃)₂-, -(CH₂)₃C(CH₃)₂-, etc.);
 (2) C₂₋₆ alkenylene (e.g. -CH=CH-, -CH₂-CH=CH-, -C(CH₃)₂-CH=CH-, -CH₂-CH=CH-CH₂-, -CH₂-CH₂-CH=CH-, -CH=CH-CH=CH-, -CH=CH-CH=CH-CH₂-, etc.);
 (3) C₂₋₆ alkynylene (e.g. -C≡C-, -CH₂-C≡C-, -CH₂-C≡C-CH₂-, etc.)

20 each of which may have 1 to 5, preferably 1 to 3, halogen atoms (e.g. fluorine, chlorine, bromine, iodine, etc.).

25 The "bivalent C₁₋₆ non-cyclic hydrocarbon groups" may have 1 to 5, preferably 1 to 3 substituents at a substitutable position. Examples of such substituents include halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), hydroxy, C₁₋₆ alkyl-carbonyloxy (e.g., acetoxy, etc.).

30 As the "bivalent C₅₋₈ monocyclic non-aromatic hydrocarbon groups", for instance, bivalent groups formed by removing an optional two hydrogen atoms from C₅₋₈ cycloalkane or C₅₋₈ cycloalkene, can be mentioned. Concrete

examples include 1,2-cyclopentylene; 1,3-cyclopentylene; 1,2-cyclohexylene; 1,3-cyclohexylene; 1,4-cyclohexylene; 1,2-cycloheptylene; 1,3-cycloheptylene; 1,4-cycloheptylene; 3-cyclohexen-1,4-ylene; 3-cyclohexen-5-1,2-ylene; 2,5-cyclohexadien-1,4-ylene. Especially, C₅₋₈ cycloalkylene is preferable.

The "spacer having a main chain of 1 to 6 atoms" is preferably a bivalent group consisting of 1 to 3 species selected from -O-, -S-, -CO-, -SO-, -SO₂-, -NR^b- (R^b has 10 the same meaning as defined above) and optionally halogenated bivalent C₁₋₆ non-cyclic hydrocarbon groups.

Preferred examples of the "spacer having a main chain of 1 to 6 atoms" include

(1) C₁₋₆ alkylene (e.g. -CH₂-, -(CH₂)₂-, -(CH₂)₃-, -(CH₂)₄-, -(CH₂)₅-, -(CH₂)₆-, -CHCH₃-, -C(CH₃)₂-, -CH(CF₃)-, -(CH(CH₃))₂-, -(CF₂)₂-, -(CH₂)₂C(CH₃)₂-, -(CH₂)₃C(CH₃)₂-, etc.);

(2) C₂₋₆ alkenylene (e.g. -CH=CH-, -CH₂-CH=CH-, -CH₂-CF=CH-, -C(CH₃)₂-CH=CH-, -CH₂-CH=CH-CH₂-, -CH₂-CH₂-CH=CH-, -CH=CH-CH=CH-, -CH=CH-CH₂-CH₂-CH₂-, etc.);

(3) C₂₋₆ alkynylene (e.g. -C≡C-, -CH₂-C≡C-, -CH₂-C≡C-CH₂-, etc.);

(4) -(CH₂)_{w1}O(CH₂)_{w2}-, -(CH₂)_{w1}S(CH₂)_{w2}-, -(CH₂)_{w1}CO(CH₂)_{w2}-, -(CH₂)_{w1}SO(CH₂)_{w2}-, -(CH₂)_{w1}SO₂(CH₂)_{w2}-, -(CH₂)_{w1}NR^b(CH₂)_{w2}-,

(5) -(CH₂)_{w3}CONR^b(CH₂)_{w4}-, -(CH₂)_{w3}NR^bCO(CH₂)_{w4}-, -(CH₂)_{w3}SO₂NR^b(CH₂)_{w4}-, -(CH₂)_{w3}NR^bSO₂(CH₂)_{w4}-, -(CH₂)_{w3}COO(CH₂)_{w4}-,

(6) -(CH₂)_{w5}NR^bCONR^b(CH₂)_{w6}-,

(7) -(CH₂)_{w7}CONR^b- (CH₂)_{w8}-CONR^b- (CH₂)_{w9}-, -CH=CH-CONR^b-; -CH=CH-SO₂NR^b-;

wherein R^b has the same meaning as defined above; R^b^b has the same meaning as R^b; w1 and w2 is an integer of 0 to 5, and w1 + w2 is 0 to 5; w3 and w4 is an integer of 0 to 4, and w3 + w4 is 0 to 4; w5 and w6 is an integer of 0 to 3, and w5 + w6 is 0 to 3; w7, w8 and w9 is an integer of

0 to 2, and w₇ + w₈ + w₉ is 0 to 2.

The "spacer having a main chain of 1 to 6 atoms" for X, is preferably -(CH₂)_{w₁}O(CH₂)_{w₂}- (symbols have the same meaning as defined above), -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}-, -SO₂NR^{8c}- (R⁸ is hydrogen atom or C₁₋₆ alkyl); more preferably -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}-, -SO₂NR^{8c}- (R⁸ has the same meaning as defined above); especially preferably -CONH-, -NHCO-, etc.

10 The "spacer having a main chain of 1 to 6 atoms" for Y, is preferably optionally halogenated bivalent C₁₋₆ non-cyclic hydrocarbon groups, -(CH₂)_{w₃}CONH(CH₂)_{w₄}-, -(CH₂)_{w₃}COO(CH₂)_{w₄}- (symbols have the same meaning as defined above); more preferably C₁₋₃ alkylene (e.g. -CH₂- , -(CH₂)₂- , -(CH₂)₃- , etc.), -(CH₂)_{w₃}CONH(CH₂)_{w₄}- , -(CH₂)_{w₃}COO(CH₂)_{w₄}- (symbols have the same meaning as defined above); especially preferably C₁₋₃ alkylene (e.g. -CH₂- , -(CH₂)₂- , -(CH₂)₃- , etc.), etc.

20 As "substituents" and "monocyclic aromatic rings" in "monocyclic aromatic rings which may be condensed with 4 to 8 membered non-aromatic rings, and may have further substituents" for Ar, those exemplified as "substituents" and "cyclic group" in the "cyclic group which may have substituents" for the above Ar¹ can be used. The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

30 The substituents are preferably formyl, optionally halogenated C₁₋₆ alkyl-carbonyl, optionally halogenated C₁₋₆ alkylsulfonyl, etc.

Here, as "optionally halogenated C₁₋₆ alkyl-carbonyl" and "optionally halogenated C₁₋₆ alkylsulfonyl", those exemplified as "substituents" in "C₇₋₁₁, aralkyl which may have substituents" can be used respectively.

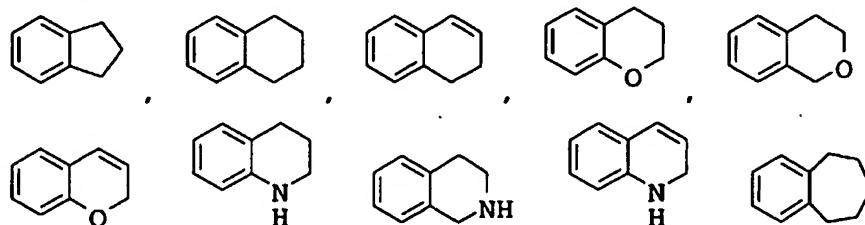
Examples of "4 to 8 membered non-aromatic rings" in the "monocyclic aromatic rings which may be condensed with 4 to 8 membered non-aromatic rings, and may have further substituents" include C₄₋₈ monocyclic non-aromatic hydrocarbon rings, 4 to 8 membered monocyclic non-aromatic hetero rings.

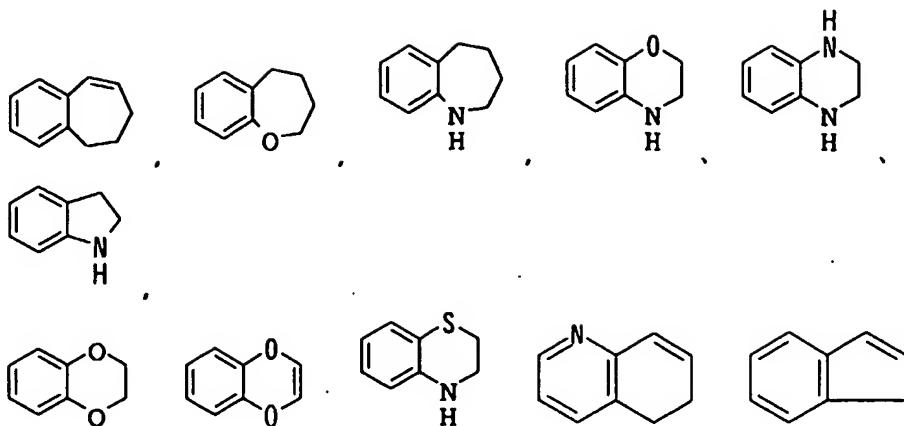
5 Examples of the "C₄₋₈ monocyclic non-aromatic hydrocarbon rings" include C₄₋₈ cycloalkane and C₄₋₈ cycloalkene. Concrete examples include cyclobutane, cyclopentane, cyclohexane, cycloheptane, cyclooctane, cyclopentene, cyclohexene, cycloheptene. Especially, cyclopentane, cyclohexane, cyclobutane, etc. are preferable.

10 Examples of the "4 to 8 membered monocyclic non-aromatic hetero rings" include azetidine, pyrrolidine, pyrroline, pyrazolidine, 2- or 3-pyrazoline, imidazoline, piperidine, piperazine, azepine, azokane, oxane, oxine, oxepane, oxazolidine, 2-oxazoline, thiazolidine, 2-thioazoline, morpholine, thiomorpholine.

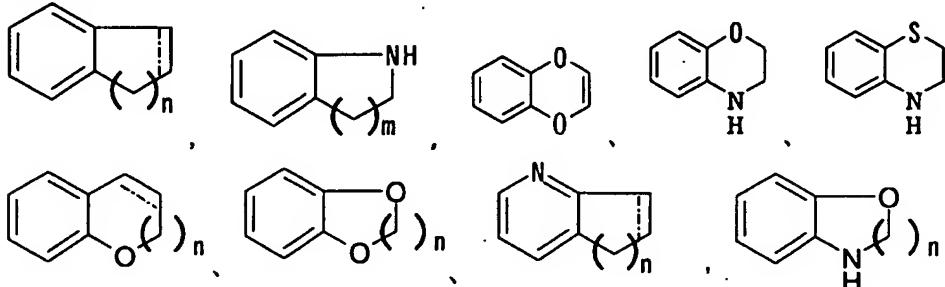
15 The above "4 to 8 membered non-aromatic rings" may have 1 to 3 substituents at a substitutable position. Examples of such substituents include optionally halogenated C₁₋₆ alkyl (e.g., methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, hexyl, etc.), cyano, hydroxy.

20 Regarding Ar, concrete examples of "monocyclic aromatic rings which may be condensed with 4 to 8 membered non-aromatic rings, and may have further substituents" include



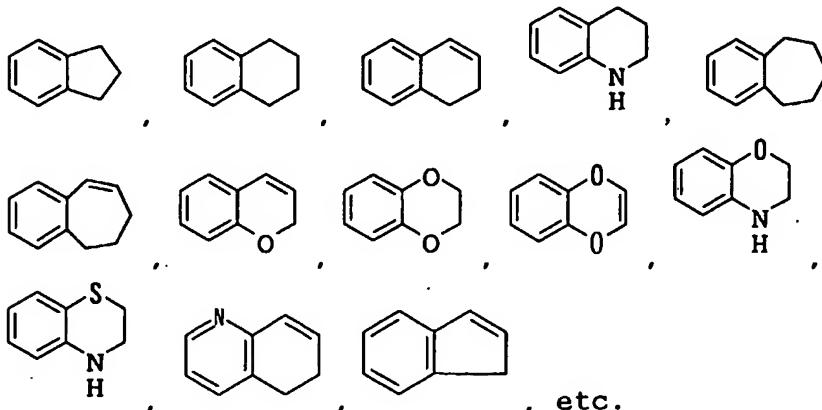


Ar is preferably benzene, pyridine, or rings of the
5 formulae :



wherein ----- is a single bond or double bond; each of m
and n is an integer of 1 to 4.

10 Ar is more preferably benzene, pyridine, rings of the
formulae :



15 As the "hydrocarbon groups which may have
substituents" for R¹ and R², those exemplified as the aboveR³

can be used.

The "hydrocarbon groups which may have substituents" are preferably "C₁₋₆ alkyl which may have substituents".

Here, examples of "C₁₋₆ alkyl" in the "C₁₋₆ alkyl which 5 may have substituents" include methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, hexyl. Especially, methyl, ethyl, propyl, etc. are preferable.

Examples of "substituents" in the "C₁₋₆ alkyl which may 10 have substituents" include halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), C₁₋₃ alkylenedioxy (e.g. methylenedioxy, ethylenedioxy etc.), nitro, cyano, optionally halogenated C₃₋₆ cycloalkyl, optionally halogenated C₁₋₆ 15 alkylthio, hydroxy, amino, mono-C₁₋₆ alkylamino (e.g. methylamino, ethylamino, propylamino, isopropylamino, butylamino, etc.), di-C₁₋₆ alkylamino (e.g. dimethylamino, diethylamino, dipropylamino, dibutylamino, ethylmethylamino, etc.), formyl, carboxy, carbamoyl, 20 thiocabamoyl, optionally halogenated C₁₋₆ alkyl-carbonyl, optionally halogenated C₁₋₆ alkoxy-carbonyl (e.g. methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, tert-butoxycarbonyl, etc.), mono-C₁₋₆ alkyl-carbamoyl (e.g. methylcarbamoyl, ethylcarbamoyl, etc.), di-C₁₋₆ alkyl- 25 carbamoyl (e.g. dimethylcarbamoyl, diethylcarbamoyl, ethylmethylcarbamoyl etc.), optionally halogenated C₁₋₆ alkylsulfonyl, formylamino, optionally halogenated C₁₋₆ alkyl-carboxamide, C₁₋₆ alkoxy-carboxamide (e.g. methoxycarboxamide, ethoxycarboxamide, 30 propoxycarboxamide, butoxycarboxamide, etc.), C₁₋₆ alkylsulfonylamino (e.g. methylsulfonylamino, ethylsulfonylamino, etc.), C₁₋₆ alkyl-carbonyloxy (e.g. acetoxy, propanoyloxy, etc.), C₁₋₆ alkoxy-carbonyloxy (e.g. methoxycarbonyloxy, ethoxycarbonyloxy, 35 propoxycarbonyloxy, butoxycarbonyloxy, etc.), mono-C₁₋₆ alkyl-carbamoyloxy (e.g. methylcarbamoyloxy,

ethylcarbamoyloxy, etc.), di-C₁₋₆ alkyl-carbamoyloxy (e.g. dimethylcarbamoyloxy, diethylcarbamoyloxy, etc.), and aromatic groups which may have substituents. The number of substituents is, for instance, 1 to 5, preferably 1 to 5. When the number of substituents is 2 or more, each substituents can be the same or different.

Here, as "optionally halogenated C₃₋₆ cycloalkyl," "optionally halogenated C₁₋₆ alkoxy" and "optionally halogenated C₁₋₆ alkylthio", those exemplified as 10 "substituents" in the above "cyclic group which may have substituents" can be used.

As "optionally halogenated C₁₋₆ alkyl-carbonyl," "optionally halogenated C₁₋₆ alkylsulfonyl" and "optionally halogenated C₁₋₆ alkyl-carboxamide", those exemplified as 15 "substituents" in the above "C₁₋₁₁, aralkyl which may have substituents" can be used.

As "substituents" and "aromatic groups" in the "aromatic groups which may have substituents", those exemplified as "substituents" and "aromatic groups" in the 20 "cyclic group which may have substituents" for the above Ar¹ can be used. The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

25 Examples of "nitrogen-containing hetero rings" in the "nitrogen-containing hetero rings which may have substituents" formed by R¹ and R² together with the adjacent nitrogen atom, include 3 to 8 membered nitrogen-containing hetero rings which contain at least one nitrogen atom in 30 addition to carbon atoms, and which may further contain 1 to 3 hetero atoms selected from nitrogen, sulfur and oxygen atom. Concrete examples include aziridine, azetidine, morpholine, thiomorpholine, piperidine, piperazine, pyrrolidine, hexamethyleneimine, heptamethyleneimine, 35 hexahydropyrimidine, 1,4-diazepan, 4,5-dihydro-imidazole, and their unsaturated cyclic amines (e.g.

1,2,5,6-tetrahydropyridine, etc.) can be mentioned. Especially, morpholine, piperidine, piperazine, pyrrolidine.

As "substituents" in the "nitrogen-containing hetero rings which may have substituents", for instance, those exemplified as "substituents" in the above "5 to 7 membered saturated cyclic amino which may have substituents" can be used. The number of substituents is, for instance, 1 to 5, preferably 1 to 3. When the number of substituents is 2 or more, each substituents can be the same or different.

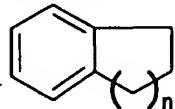
R¹ and R² are preferably C₁₋₆ alkyl, more preferably methyl, ethyl, propyl, etc.

Also, it is preferable that R¹ and R², together with the adjacent nitrogen atom, form piperidino, 15 pyrrolidin-1-yl, piperazin-1-yl etc.

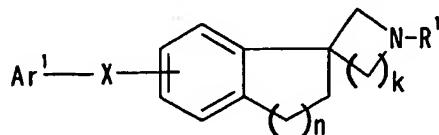
And, it is preferable that at least one of R¹ and R² is C₁₋₆ alkyl which may have substituents. It is especially preferable that both R¹ and R² is C₁₋₆ alkyls which may have substituents.

20

R² can form a spiro ring together with Ar. For instance, Ar is a ring of the formula :



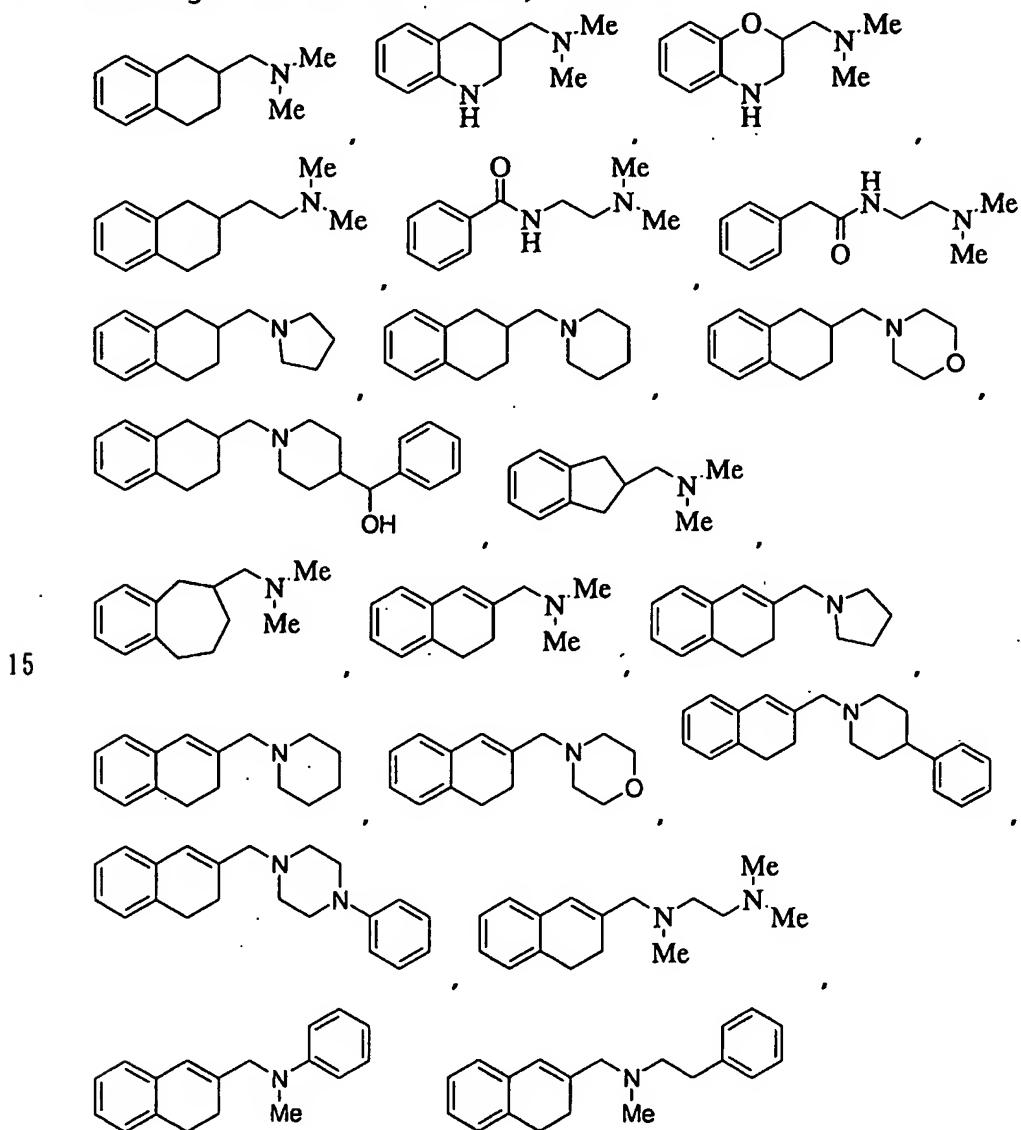
wherein n is an integer of 1 to 4; and Y is methylene; R² can form a spiro ring together with Ar. Examples of the spiro ring include

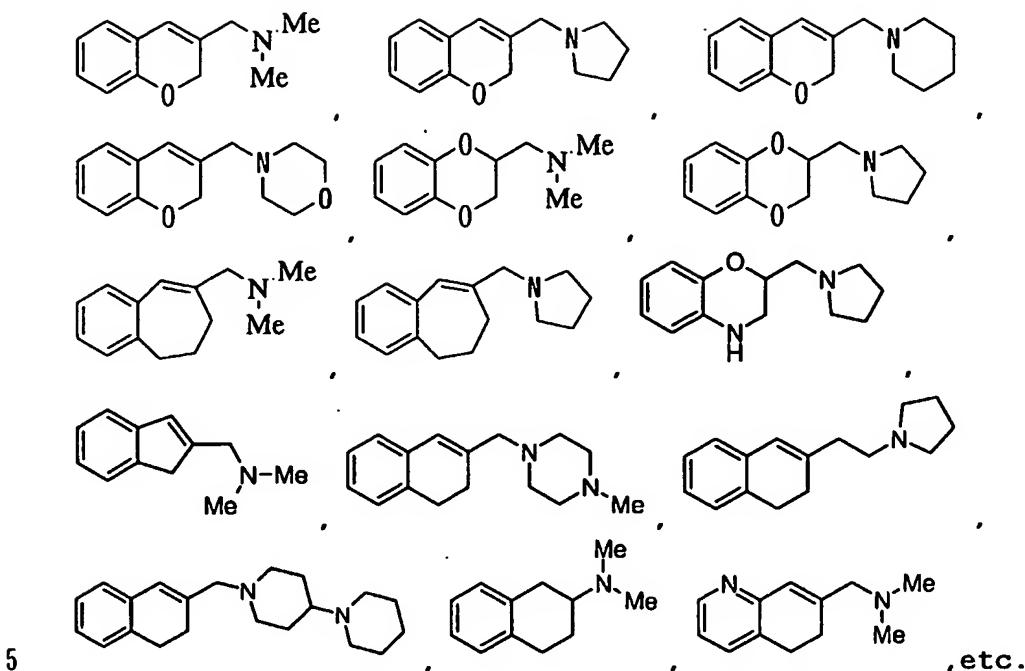


wherein k (ring Ar and N are connected by -(CH₂)_k-) is an integer of 1 to 4; and other symbols have the same meaning 30 as defined above.

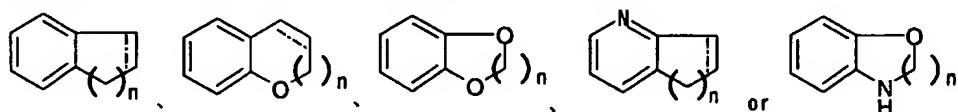
R^2 may form, together with the adjacent nitrogen atom and Y, a nitrogen-containing hetero ring which may have substituents. Examples of the "nitrogen-containing hetero ring which may have substituents" include those exemplified as the "nitrogen-containing hetero rings which may have substituents" formed by R^1 and R^2 together with the adjacent nitrogen atom.

In formula (I), preferable examples of the partial structural formula : Ar-Y-N(R^1) R^2 (symbols have the same meanings as defined above) include



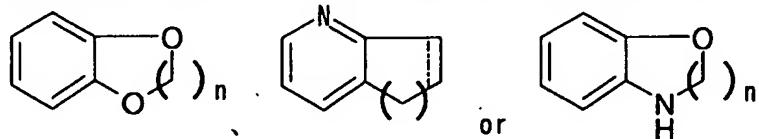


Among the compounds of the formula (I), a compound wherein Ar is a ring of the formula :



wherein ----- is a single bond or double bond, n is an integer of 1 to 4, and each ring may have substituents;
 10 X is $\text{-CONR}^{\text{8c}}\text{-}$, $\text{-NR}^{\text{8c}}\text{CO-}$, $\text{-CH=CH-CNR}^{\text{8c}}\text{-}$ or $\text{-SO}_2\text{NR}^{\text{8c}}\text{-}$ where R⁸ is hydrogen atom or C₁₋₆ alkyl;
 Y is a spacer having a main chain of 1 to 6 atoms;

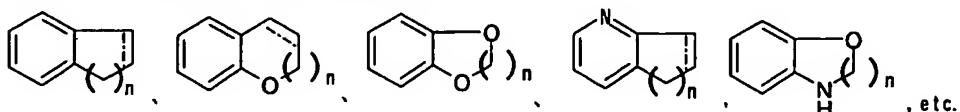
provided that Ar is a ring of the formulae :



15 wherein symbols have the same meanings as defined above, and each ring may have substituents, when X is $\text{-SO}_2\text{NH-}$; and provided that Ar¹ is not biphenylyl which may be substituted; when X is -CONH- and Ar is any one of 20 benzopyran, dihydrobenzopyran, dihyrobenzoxazine, dihydrobenzoxazole or tetrahydrobenzoxazepine;

(excluding N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-4-biphenylylcarboxamide);
 namely compound of the formula (I') (excluding N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-4-biphenylylcarboxamide) is a novel compound.

Preferred examples of compound of the formula (I') include compound of the formula (I'-1), (I'-2), (I'-3), (I'-4), (I'-5), (I'-6), (I'-7), (I'-8), (I'-9) or (I'-10).
 In the above formulae (I'), (I'-1), (I'-2), (I'-3), (I'-4), (I'-5), (I'-6), (I'-7), (I'-8), (I'-9) and (I'-10), a ring of the formula :



wherein symbols have the same meanings as above,
 may have further 1 to 3 substituents at substitutable
 15 positions.

Examples of such substituents include "substituents" exemplified in the above Ar. Especially, preferred are formyl, optionally halogenated C₁₋₆ alkyl-carbonyl, optionally halogenated C₁₋₆ alkylsulfonyl, optionally 20 halogenated C₁₋₆ alkyl (e.g., methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, tert-butyl, pentyl, hexyl, etc.), cyano, hydroxy, etc.

Examples of salts of compound (I) or (I') include salts 25 with inorganic bases, ammonium salts, salts with organic bases, salts with inorganic acids, salts with organic acids, and salts with basic or acidic amino acids.

Preferred examples of salts with inorganic bases include alkali metal salts such as sodium salts and 30 potassium salts; alkaline earth metal salts such as calcium salts, magnesium salts, barium salts; and aluminum salts.

Preferred examples of salts with organic bases include salts with trimethylamine, triethylamine, pyridine, picoline, ethanolamine, diethanolamine, triethanolamine,

dicyclohexylamine, N,N-dibenzylethylenediamine.

Preferred examples of salts with inorganic acids include salts with hydrochloric acid, hydrobromic acid, nitric acid, sulfuric acid, phosphoric acid.

5 Preferred examples of salts with organic acids include salts with formic acid, acetic acid, trifluoroacetic acid, fumaric acid, oxalic acid, tartaric acid, maleic acid, citric acid, succinic acid, malic acid, methanesulfonic acid, benzenesulfonic acid, and p-toluenesulfonic acid,

10 3-chlorobenzoic acid.

Preferred examples of salts with basic amino acids include salts with arginine, lysine, ornithine. Preferred examples of salts with acidic amino acids include salts with aspartic acid, glutamic.

15 Among these salts, pharmaceutically acceptable salts are preferable. For instance, when compound (I) or (I') possesses an acidic functional group, it can form an inorganic salt such as an alkali metal salt (e.g., sodium salt, potassium salt, etc.), an alkaline earth metal salt (e.g. calcium salt, magnesium salt, barium salt, etc.), and an ammonium salt. When compound (I) or (I') possesses a basic functional group, it can form an inorganic salt such as hydrochloride, sulfate, phosphate, hydrobromate, etc.; or an organic salt such as acetate, maleate, fumarate, 20 succinate, methanesulfonate, p-toluenesulfonate, citrate and tartrate.

25 Compounds (I) and (I') (hereinafter also abbreviated as a compound of the invention) can be either anhydrides or hydrates. A hydrate may have 0.5 to 3 water molecules.

30 In addition, a compound of the invention can be labeled using isotopes (e.g. ³H, ¹⁴C, and ³⁵S, etc.).

When a compound of the invention contains optical isomers, stereoisomers, regio isomers, rotational isomers, 35 these are included as a compound of the invention, and each of them can be obtained as a single substance by per se known

synthesis methods and separation methods. For instance, when optical isomers exist in a compound of the invention, the optical isomers separated from the compound are included in a compound of the invention.

5 The optical isomers can be produced using per se known methods. Concretely, the optical isomer can be obtained by using an optically active synthetic intermediate, or subjecting the racemic mixture of the final product to optical resolution in accordance with common method.

10 Examples of optical resolution methods include per se known methods such as the fractional recrystallization method, chiral column method, diastereomer method, etc., which are described in detail below.

1) Fractional recrystallization method

15 The method which comprises allowing a racemate to form a salt with an optically active compound (e.g. (+)-mandelic acid, (-)-mandelic acid, (+)-tartaric acid, (-)-tartaric acid, (+)-1-phenethylamine, (-)-1-phenethylamine, cinchonine, (-)-cinchonidine, brucine, etc.), separating 20 the salt using a fractional recrystallization method, followed by, if desired, neutralizing process to obtain a free optical isomer.

2) Chiral column method

25 This method comprises subjecting a racemate or its salt to a column for separating an optical isomer (chiral column) for separation. For instance, in the case of liquid chromatography, an optical isomer mixture is added to the chiral column such as ENANTIO-OVM [produced by Toso] or CHIRAL series [produced by Daicel], which is developed 30 using water, various buffer solutions (e.g. phosphate buffer), organic solvents (e.g. ethanol, methanol, isopropanol, acetonitrile, trifluoroacetic acid, diethylamine, etc.) as single or mixed solutions, and the optical isomers are separated. Also, in the case of gas 35 chromatography, for instance, separation is conducted using a chiral column such as CP-Chirasil-DeX (produced by

G.L.Science Co.).

3) Diastereomer method

In this method, a racemic mixture is subjected to a chemical reaction with an optically active reagent to give 5 a diastereomer mixture, which is separated into a single substance by an ordinary separation means (e.g. fractional recrystallization, chromatography method, etc.). This single substance is subjecting to removal of the optically active reagent part using chemical processing such as a 10 hydrolysis reaction. For instance, when a compound of the invention possesses hydroxy or primary or secondary amino in its molecule, this compound is subjected to a condensation reaction with an optically active organic acid (e.g. MTPA [α -methoxy- α -(trifluoromethyl)phenylacetic acid], (-)-menthoxyacetic acid, etc.), to give the 15 diastereomer in an ester form or an amide form, respectively. On the other hand, when a compound of the invention possesses carboxylic acid group, this compound is subjected to a condensation reaction with an optically 20 active amine or alcohol reagent, to give the diastereomer in an amide form or an ester form, respectively. The separated diastereomer can be converted to an optical isomer of the original compound, by applying acidic 25 hydrolysis or basic hydrolysis.

A prodrug of compound (I') is a compound which is converted to compound (I') by reactions involving enzymes and gastric acid, etc. under physiological conditions in the living body; in other words, a compound that is changed into compound (I') by enzymatically-caused oxidation, 30 reduction and hydrolysis, and a compound that is changed into compound (I') by hydrolysis caused by gastric acid. Examples of the prodrugs of compound (I') include compounds in which amino groups of compound (I') have been acylated, alkylated, or phosphorylated [e.g. compounds in 35 which amino groups of compound (I') have been eicosanoylated, aranylated, pentylaminocarbonylated,

(5-methyl-2-oxo-1,3-dioxolen-4-yl)methoxycarbonylated, tetrahydrofuranylated, pyrrolidylmethylated, pivaloyloxymethylated, tert-butylated, etc.]; compounds in which hydroxyl groups of compound (I') have been 5 acylated, alkylated, phosphorylated, borated (e.g. compounds in which hydroxyl groups of compound (I') have been acetylated, palmitoylated, propanoylated, pivaloylated, succinylated, fumarilated, alanilated, dimethylaminomethylcarbonylated, etc.); compounds in 10 which carboxyl groups of compound (I') have been esterified or amidated [e.g. compounds in which carboxyl groups of compound (I') have been ethylesterified, phenylesterified, carboxylmethylesterified, dimethylaminomethylesterified, 15 pivaloyloxymethylesterified, ethoxycarbonyloxyethylesterified, phthalidylesterified, (5-methyl-2-oxo-1,3-dioxolen-4-yl)methylesterified, cyclohexyloxycarbonylethylesterified, or methylamidated, etc.]. These compounds can be produced from compound (I') 20 using per se known methods.

Also, a prodrug of compound (I') can be a compound which is changed to compound (I') by physiological conditions, as described in pages 163 to 198 of Molecular Design, Volume 7, "Development of Drugs.", published in 25 1990 by Hirokawa Shoten.

A compound of the invention can be produced in accordance with per se known methods such as methods described in WO9838156, WO9532967, and EP-A533266, etc., 30 or analogous methods thereto.

For instance, a compound of the invention can be produced in accordance with [Production method 1] to [Production method 6] which are described in detail below, or analogous methods thereto.

35 Compounds (II) to (XI) used as raw materials, can be used in the form of salts. As such salts, those exemplified

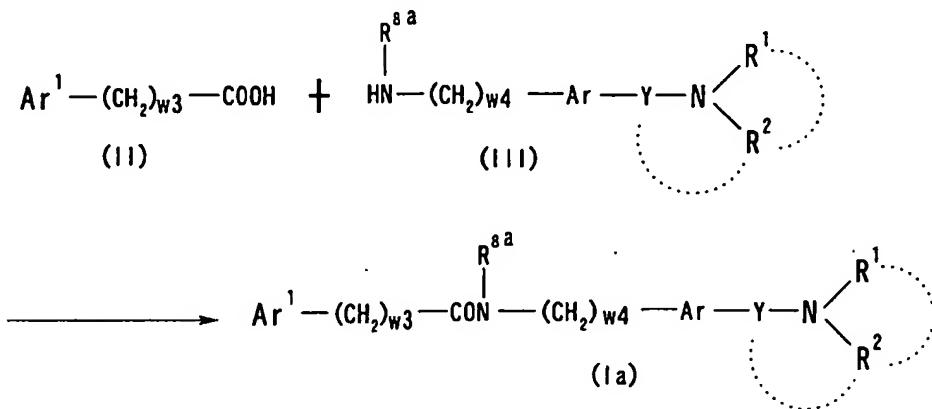
as salts of the above compound (I) or (I') can be used.

In the following [Production method 1] to [Production method 6], when an alkylation reaction, a hydrolysis reaction, an amination reaction, an esterification reaction, an amidation reaction, an esterification reaction, an etherification reaction, an oxidation reaction, a reduction reaction, etc. are carried out, these reactions are carried out in accordance with per se known methods. Examples of such methods include the methods described in *Organic Functional Group Preparations*, Second Edition, Academic Press, Inc., published in 1989; *Comprehensive Organic Transformations*, VCH Publishers Inc., published in 1989, etc.

15 [Production method 1]

Compound (Ia) having $-(\text{CH}_2)_{w_3}\text{CONR}^{8a}(\text{CH}_2)_{w_4}-$ for X in formula (I), is produced, for instance, by the following amidation reaction.

(Amidation reaction)



20 wherein R^{8a} is hydrogen atom or an optionally halogenated C_{1-6} alkyl; other symbols have the same meanings as defined above.

As the "optionally halogenated C_{1-6} alkyl", those exemplified as "substituents" in the above "cyclic group which may have substituents" can be used.

The "amidation reaction" includes the following

"method using a dehydration and condensation agent" and "method using a reactive derivative of carboxylic acid".

i) Method using a dehydration and condensation agent

5 Compound (III), 1 to 5 equivalents of compound (II), and 1 to 2 equivalents of a dehydration and condensation agent are reacted in an inert solvent. If necessary, the reaction can be carried out with the coexistence of 1 to 1.5 equivalents of 1-hydroxybenzotriazole (HOBT) and (or)

10 catalytic quantity to 5 equivalents of a base.

Examples of the "dehydrating and condensation agent" include dicyclohexylcarbodiimide (DCC), 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride (WSC). WSC is particularly preferable.

15 Examples of the "inert solvent" include nitrile solvents (preferably acetonitrile), amide solvents (preferably DMF), halogenated hydrocarbon solvents (preferably dichloromethane), ether solvents (preferably THF). Two or more kinds of these can be mixed in an

20 appropriate ratio for use.

Examples of the "base" include

1) for instance, strong bases such as hydrides of alkali metals or alkaline earth metals (e.g. lithium 25 hydride, sodium hydride, potassium hydride, calcium hydride, etc.), amides of alkali metals or alkaline earth metals (e.g. lithium amide, sodium amide, lithium diisopropylamide, lithium dicyclohexylamide, lithium hexamethyldisilazide, sodium hexamethyldisilazide, 30 potassium hexamethyldisilazide, etc.), lower alkoxides of alkali metals or alkaline earth metals (e.g. sodium methoxide, sodium ethoxide, potassium tert-butoxide, etc.);

2) for instance, inorganic bases such as hydroxides 35 of alkali metals or alkaline earth metals (e.g. sodium hydroxide, potassium hydroxide, lithium hydroxide, barium

hydroxide, etc.), carbonates of alkali metals or alkaline earth metals (e.g. sodium carbonate, potassium carbonate, cesium carbonate, etc.) and hydrogencarbonates of alkali metals or alkaline earth metals (e.g. sodium 5 hydrogencarbonate, potassium hydrogencarbonate, etc.); and

3) for instance, amines such as triethylamine, diisopropylethylamine, N-methylmorpholine, dimethylaminopyridine, DBU (1,8-10 diazabicyclo[5.4.0]undec-7-en), DBN (1,5-diazabicyclo[4.3.0]non-5-en); for instance, organic bases such as basic heterocyclic compounds of pyridine, imidazole, 2,6-lutidine, etc.

Among the above bases, triethylamine, 4-dimethylaminopyridine, etc., are preferable.

Reaction temperature is usually room temperature (0°C to 30°C, hereafter the same). Reaction time is, for instance, 10 to 24 hours.

ii) Method using a reactive derivative of carboxylic acid
A reactive derivative of compound (II) and 1 to 5 equivalents (preferably 1 to 3 equivalents) of compound (III) are reacted in an inert solvent. If necessary, the reaction can be carried out with the coexistence of 1 to 10 equivalents, preferably 1 to 3 equivalents of a base.

Examples of the "reactive derivative" of compound (II) include acid halides (e.g., acid chloride, acid bromide, etc.), mixed acid anhydrides (e.g. acid anhydrides with C₁₋₆ alkyl-carboxylic acid, C₆₋₁₀ aryl-carboxylic acid or C₁₋₆ alkylcarbonate), active esters (e.g. esters with phenol 30 which may have substituents, 1-hydroxybenzotriazole or N-hydroxysuccinimide, etc.).

Examples of the "substituents" in the "phenol which may have substituents" include halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), nitro, optionally 35 halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkoxy.

The number of substituents is, for instance, 1 to 5.

As the "optionally halogenated C₁₋₆ alkyl" and "optionally halogenated C₁₋₆ alkoxy", those exemplified as "substituents" in the above "cyclic group which may have substituents" can be used.

5 Concrete examples of "phenol which may have substituents" include phenol, pentachlorophenol, pentafluorophenol, p-nitrophenol. The reactive derivative is, preferably, an acid halide.

Examples of the "inert solvent" include ether solvents, halogenated hydrocarbon solvents, aromatic solvents, nitrile solvents, amide solvents, ketone solvents, sulfoxide solvents, and water. Two or more kinds of these can be mixed in an appropriate ratio for use. Especially, acetonitrile, THF, dichloromethane, chloroform, etc. are preferable.

As the "base", the same as above are used. The base is preferably sodium hydride, potassium carbonate, sodium carbonate, sodium hydroxide, potassium hydroxide, sodium hydrogencarbonate, potassium hydrogencarbonate, triethylamine, pyridine, etc.

Reaction temperature is usually -20°C to 50°C, preferably room temperature. Reaction time is usually 5 minutes to 40 hours, preferably 1 to 18 hours.

Compound (III) can be produced by per se known methods. For instance, 6-amino-2-(N,N-dimethylamino)methyltetraline or its salt can be produced in accordance with the methods described in WO9838156. Also, 6-amino-2,3-dihydro-1-(2-dimethylaminoethyl)-1H-indole, 6-amino-3,4-dihydro-4-(2-dimethylaminoethyl)-2H-1,4-benzoxazine, etc., can be produced in accordance with the methods described in WO9532967.

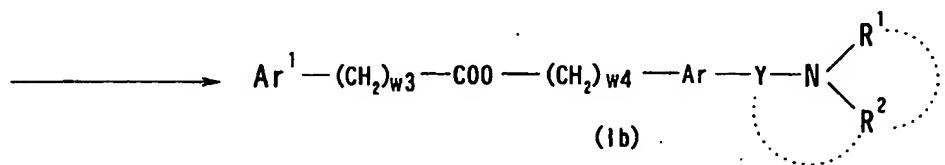
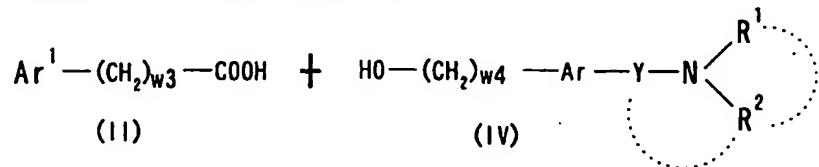
The above "method using a reactive derivative of carboxylic acid" can be also adopted when producing a corresponding sulfonamide derivative or sulfinamide derivative, from the sulfonic acid of the formula : Ar¹-(CH₂)_n-SO₃OH (symbols have the same meanings as defined

above), or the sulfinic acid of the formula : $\text{Ar}^1-(\text{CH}_2)_w-\text{SOOH}$ (symbols have the same meanings as defined above).

[Production method 2]

5 Compound (Ib) having $-(CH_2)_{w_3}-COO(CH_2)_{w_4}-$ for X in the formula (I), can be produced by the following esterification reaction.

(Esterification reaction)



10 wherein symbols have the same meanings as defined above.

A reactive derivative of compound (II) and 1 to 5 equivalents (preferably 1 to 3 equivalents) of compound (IV) is reacted in an inert solvent. Usually, this reaction is carried out with the coexistence of 1 to 10 equivalents, preferably 1 to 3 equivalents of a base.

As the reactive derivative of compound (II), the same as above is used. Especially, an acid halide is preferable.

Examples of the "inert solvent" include ether solvents, halogenated hydrocarbon solvents, aromatic solvents, nitrile solvents, amide solvents, ketone solvents, sulfoxide solvents. Two or more kinds of these can be mixed in an appropriate ratio for use. Especially, acetonitrile, dichloromethane, chloroform, etc. are preferable.

25 As the "base", the same one as above can be used. The base is preferably sodium hydride, potassium carbonate, sodium carbonate, sodium hydroxide, potassium hydroxide, sodium hydrogencarbonate, potassium hydrogencarbonate,

triethylamine, pyridine, etc.

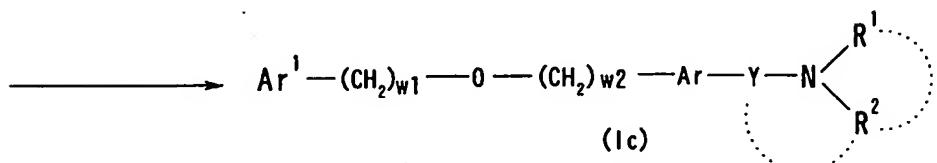
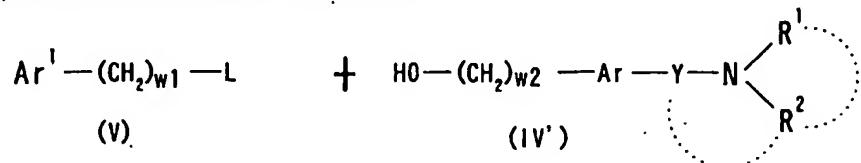
Reaction temperature is usually -20°C to 50°C, preferably room temperature. Reaction time is usually 5 minutes to 40 hours, preferably 1 to 18 hours.

5

[Production method 3]

Compound (Ic) having $-(CH_2)_{w1}O(CH_2)_{w2}-$ for Y in the formula (I), can be produced by, for instance, the following etherification reaction.

10 (Etherification reaction)



wherein L is a leaving group, and other symbols have the same meanings as defined above.

Examples of the "leaving group" for L include halogen atom (e.g. chlorine, bromine, iodine, etc.), optionally halogenated C₁₋₆ alkylsulfonyloxy (e.g. methanesulfonyloxy, ethanesulfonyloxy, trifluoromethanesulfonyloxy, etc.), C₆₋₁₀ arylsulfonyloxy which may have substituents, hydroxy.

20 Examples of the "substituents" in the "C₆₋₁₀ arylsulfonyloxy which may have substituents" include halogen atom (e.g. chlorine, bromine, iodine, etc.), optionally halogenated C₁₋₆ alkyl, C₁₋₆ alkoxy. The number of substituents is, for instance, 1 to 3. Concrete examples 25 of the C₆₋₁₀ arylsulfonyloxy which may have substituents" include benzenesulfonyloxy, p-toluenesulfonyloxy, 1-naphthalenesulfonyloxy, 2-naphthalenesulfonyloxy.

The "leaving group" is preferably halogen atom (e.g.

chlorine, bromine, iodine, etc.), methanesulfonyloxy, trifluoromethanesulfonyloxy, p-toluenesulfonyloxy.

Compound (IV') and about 1 to 5 equivalents
5 (preferably 1 to 2 equivalents) of compound (V) are reacted
in inert solvent, with the coexistence of base.

As the "base", the same one as above can be used. The
base is preferably potassium carbonate, sodium
hydrogencarbonate, triethylamine, N-methylmorpholine,
10 pyridine, etc. The amount of the base used is usually about
1 to 5 equivalents relative to compound (V).

Examples of the "inert solvent" include alcohol
solvents, ether solvents, halogenated hydrocarbon
solvents, aromatic solvents, nitrile solvents, amide
15 solvents, ketone solvents, sulfoxide solvents, water. Two
or more kinds of these can be mixed in an appropriate ratio
for use. Especially, acetonitrile, N,N-dimethylformamide
(DMF), acetone, ethanol, pyridine, etc., are preferable.

Reaction temperature is about -20°C to 100°C,
20 preferably room temperature to 80°C. Reaction time is, for
instance, 5 hours to 1 day.

In the above production method, when the leaving group
is hydroxy, Mitsunobu reaction can usually be used. In the
Mitsunobu reaction, compound (V) and 0.5 to 5 equivalents
25 (preferably 1 to 1.5 equivalents) of compound (IV') are
reacted in inert solvent with the coexistence of 0.5 to 5
equivalents (preferably 1 to 1.5 equivalents) of ethyl
acetyldicarboxylate.

Examples of the inert solvent include ether solvents,
30 halogenated hydrocarbon solvents, aromatic solvents,
nitrile solvents, amide solvents, ketone solvents,
sulfoxide solvents. Two or more kinds of these can be mixed
in an appropriate ratio for use. Especially,
acetonitrile, dichloromethane, chloroform, etc. are
35 preferable.

Reaction temperature is usually -20°C to 50°C.

preferably room temperature. Reaction time is usually 5 minutes to 40 hours, preferably 1 to 18 hours.

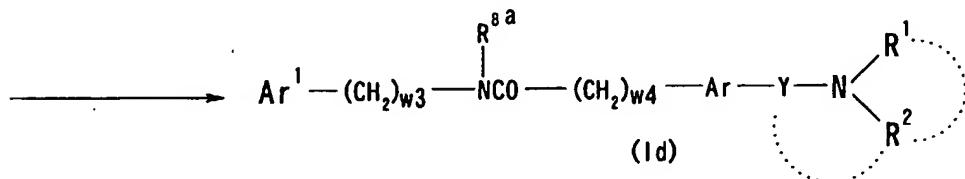
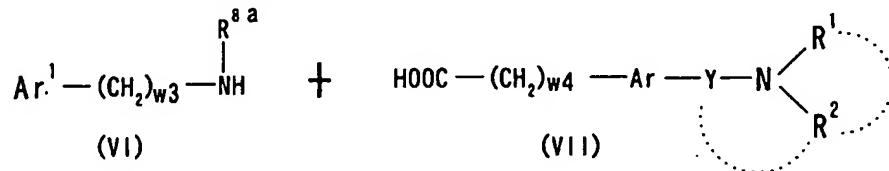
Compound (IV') can be produced by per se known methods.

For instance, 3-(N,N-dimethylamino)methyl-1,2,3,4-tetrahydro-7-quinolinol, 2-(N,N-dimethylamino)methyl-6-hydroxytetralin, 6-hydroxy-2-piperidinomethyltetralin, 2-[2-(N,N-dimethylamino)ethyl]-6-hydroxytetralin, 2-(N,N-dimethylamino)methyl-7-hydroxytetralin, 6-hydroxy-2-(N-methylamino)methyltetralin, etc., can be produced in accordance with the methods described in WO9838156.

[Production method 4]

Compound (Id) having $-(\text{CH}_2)_{w_3}\text{NR}^{8a}\text{CO}(\text{CN}_2)_{w_4}-$ for X in the formula (I), can be produced, for instance, by the following amidation reaction.

(Amidation reaction)



wherein symbols have the same meanings as defined above.

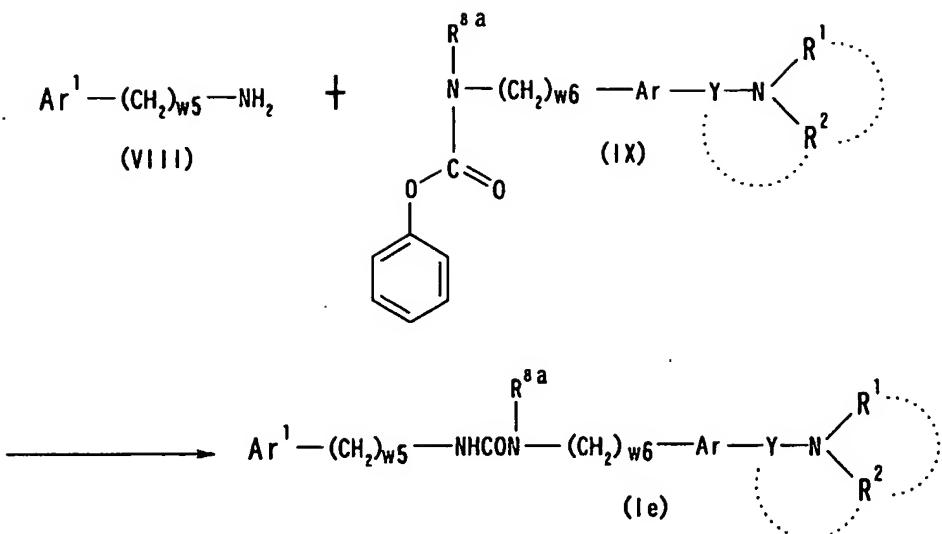
This Production method is carried out in accordance
20 with the above Production method 1.

[Production method 5]

Compound (Ie) having $-(\text{CH}_2)_{w_5}\text{NHCONR}^{8a}(\text{CN}_2)_{w_6}-$ for X in the formula (I), can be produced, for instance, by the following urea reaction.

(Urea reaction)

70



wherein symbols have the same meanings as defined above.

Compound (IX) and 1 to 5 equivalents (preferably 1 to
5 equivalents) of compound (VIII) is reacted in an inert
solvent with the coexistence of a base.

As the "base", the same one as above can be used. The
base is preferably potassium carbonate, sodium carbonate,
sodium hydroxide, potassium hydroxide, sodium
10 hydrogencarbonate, potassium hydrogencarbonate,
triethylamine, pyridine, etc.

Examples of the "inert solvent" include alcohol
solvents, ether solvents, halogenated hydrocarbon
solvents, aromatic solvents, nitrile solvents, amide
15 solvents, ketone solvents, sulfoxide solvents, water. Two
or more kinds of these can be mixed in an appropriate ratio
for use. Especially, acetonitrile, DMF, acetone, ethanol,
pyridine, etc. are preferable.

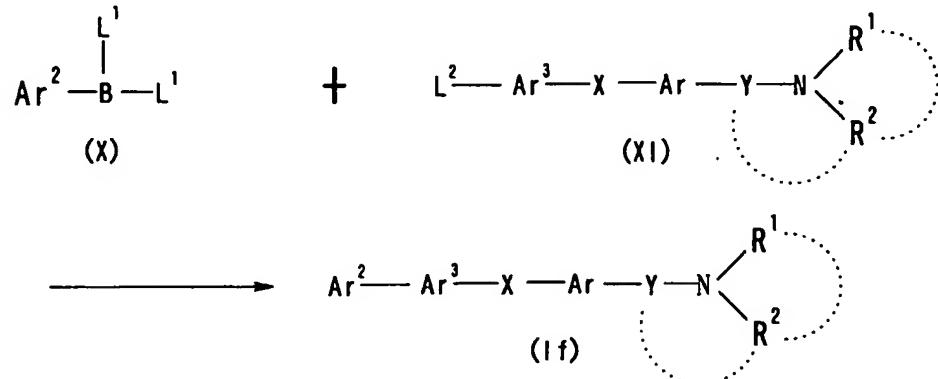
Reaction temperature is usually -20°C to 100°C,
20 preferably room temperature to 80°C. Reaction time is, for
instance, 0.5 hour to 1 day.

[Production method 6]

Compound (If) having, for Ar¹, a ring assembly aromatic

group ($\text{Ar}^2\text{-Ar}^3$) which may have substituents in the formula (I), can be produced by, for instance, the following aryl-coupling reaction.

(Aryl-coupling reaction)



5

wherein Ar² and Ar³ are monocyclic aromatic groups or condensed aromatic groups, each of which may have substituents; L¹ is hydroxy or C₁₋₆ alkyl; L² is halogen (preferably chlorine, bromine) or

10 trifluoromethanesulfonyloxy; other symbols have the same meanings as defined above.

As "substituents", "monocyclic aromatic groups" and "condensed aromatic groups" in the "monocyclic aromatic groups or condensed aromatic groups, each of which may have substituents" for Ar^2 and Ar^3 , those exemplified as the above Ar^1 can be used. Especially, it is preferable that both of Ar^2 and Ar^3 are phenyl groups which may have substituents, and $\text{Ar}^2\text{-}\text{Ar}^3$ is biphenylyl which may have substituents.

20 The aryl-coupling reaction can be carried out in accordance with per se known methods such as the method described in Acta. Chemica Scandinavia, pp. 221-230, 1993, or methods analogous thereto.

Compound (X) and 1 to 3 equivalents (preferably 1 to 1.5 equivalents) of compound (XI) are reacted in an inert solvent in the presence of a base and a transition metal catalyst.

As the base, the same one as above can be used. The

base is preferably sodium carbonate, sodium hydrogencarbonate, etc.

The amount of the "base" used is, for instance, about 1 to 10 equivalents relative to compound (XI).

5 Examples of the "transition metal catalyst" include palladium catalyst, nickel catalyst. Examples of the "palladium catalyst" include tetrakis(triphenylphosphine)palladium (0), palladium acetate, bis (triphenylphosphine) palladium (II) chloride, 10 palladium-carbon. Examples of the "nickel catalyst" include tetrakis(triphenylphosphine) nickel (0).

The amount of the "transition metal catalyst" used is about 0.01 to 1 equivalent, preferably about 0.01 to 0.5 equivalent, relative to compound (XI).

15 Reaction temperature is room temperature to 150°C, preferably about 80°C to 150°C. Reaction time is, for instance, about 1 to 48 hours.

20 Examples of the "inert solvent" include water, alcohol solvents, aromatic solvents. Two or more kinds of these can be mixed in an appropriate ratio for use. Especially, a single solvent such as water, ethanol and toluene; or a mixed solvent of two or more kinds of these is preferable.

Examples of the above "alcohol solvents" include methanol, ethanol, isopropanol, tert-butanol.

25 Examples of the above "ether solvents" include diethylether, tetrahydrofuran (THF), 1,4-dioxane, 1,2-dimethoxyethane.

30 Examples of the above "halogenated hydrocarbon solvents" include dichloromethane, chloroform, 1,2-dichloroethane, carbon tetrachloride.

Examples of the above "aromatic solvents" include benzene, toluene, xylene, pyridine.

Examples of the above "hydrocarbon solvents" include hexane, pentane, cyclohexane.

35 Examples of the above "amide solvents" include N,N-dimethylformamide (DMF), N,N-dimethylacetamide, N-

methylpyrrolidone.

Examples of the above "ketone solventd" include acetone, methylethylketone.

Examples of the above "sulfoxide solvents" include 5 dimethylsulfoxide (DMSO).

Examples of the above "nitrile solvents" include acetonitrile, propionitrile.

In a compound of the invention thus obtained, the intramolecular functional group can be converted to a 10 desired functional group by combining per se known chemical reactions. Examples of the chemical reactions include oxidation reaction, reduction reaction, alkylation reaction, hydrolysis reaction, amination reaction, esterification reaction, aryl-coupling reaction, 15 deprotection reaction.

In each of the above reactions, when the raw material compounds possess amino, carboxy, hydroxy, and/or carbonyl as substituents, protecting groups which are generally used in peptide chemicals, etc., can be introduced into these 20 groups, and the desired compound can be obtained by removing the protecting groups after the reaction if necessary.

Examples of the protecting group for amino include formyl, C₁₋₆ alkyl-carbonyl (e.g. acetyl, propionyl, etc.), C₁₋₆ alkoxy-carbonyl (e.g. methoxycarbonyl, 25 ethoxycarbonyl, tert-butoxycarbonyl, etc.), benzoyl, C₇₋₁₀ aralkyl-carbonyl (e.g. benzylcarbonyl, etc.), C₇₋₁₄ aralkyloxy-carbonyl (e.g. benzyloxycarbonyl, 9-fluorenylmethoxycarbonyl, etc.), trityl, phthaloyl, N,N-dimethylaminomethylene, silyl (e.g. trimethylsilyl, 30 triethylsilyl, dimethylphenylsilyl, tert-butyldimethylsilyl, tert-butyldiethylsilyl, etc.), C₂₋₆ alkenyl (e.g. 1-allyl, etc.). These groups may be substituted by 1 to 3 of halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), C₁₋₆ alkoxy (e.g. methoxy, 35 ethoxy, propoxy, etc.) or nitro, etc.

Examples of the protecting group for carboxy include

C_{1-6} alkyl (e.g. methyl, ethyl, propyl, isopropyl, butyl, tert-butyl, etc.), C_{7-11} aralkyl (e.g. benzyl, etc.), phenyl, trityl, silyl (e.g. trimethylsilyl, triethylsilyl, dimethylphenylsilyl, tert-butyldimethylsilyl, tert-
5 butyldiethylsilyl, etc.), C_{2-6} alkenyl (e.g. 1-allyl, etc.). These groups may be substituted by 1 to 3 of halogen atom (e.g. fluorine, chlorine, bromine, iodine, etc.), C_{1-6} alkoxy (e.g. methoxy, ethoxy, propoxy, etc.) or nitro.

Examples of the protective group for hydroxy include
10 C_{1-6} alkyl (e.g. methyl, ethyl, propyl, isopropyl, butyl, tert-butyl, etc.), phenyl, trityl, C_{7-10} aralkyl (e.g. benzyl, etc.), formyl, C_{1-6} alkyl-carbonyl (e.g. acetyl, propionyl, etc.), benzoyl, C_{7-10} aralkyl-carbonyl (e.g. benzylcarbonyl, etc.), 2-tetrahydropyranyl, 2-
15 tetrahydrofuranyl, silyl (e.g. trimethylsilyl, triethylsilyl, dimethylphenylsilyl, tert-butyldimethylsilyl, tert-butyldiethylsilyl, etc.), C_{2-6} alkenyl (e.g. 1-allyl, etc.). These groups may be substituted by 1 to 3 of halogen atom (e.g. fluorine,
20 chlorine, bromine, iodine, etc.), C_{1-6} alkyl (e.g. methyl, ethyl, n-propyl, etc.), C_{1-6} alkoxy (e.g. methoxy, ethoxy, propoxy, etc.) or nitro, etc. can be substituted for these groups.

Examples of the protecting group for carbonyl include
25 cyclic acetal (e.g. 1,3-dioxane, etc.), and non-cyclic acetal (e.g. di- C_{1-6} alkylacetal, etc.).

Removal of the above protecting groups can be carried out in accordance with per se known methods such as those described in Protective Groups in Organic Synthesis,
30 published by John Wiley and Sons (1980). For instance, the methods using acid, base, ultraviolet light, hydrazine, phenylhydrazine, sodium N-methyldithiocarbamate, tetrabutylammonium fluoride, palladium acetate, trialkylsilyl halide (e.g. trimethylsilyl iodide, trimethylsilyl bromide, etc.), and a reduction method, etc.
35 can be used.

A compound of the invention can be isolated and purified by per se known methods such as solvent extraction, changing of liquid properties, transdissolution, crystallization, recrystallization, chromatography, etc.

5 It is also possible to isolate and purify the raw material compounds of a compound of the invention, or their salts using the same known methods as above, but they can also be used as raw materials in the next process as a reaction mixture without being isolated.

10

A compound of the invention possesses an excellent MCH receptor antagonistic action, therefore, it is useful as an agent for preventing or treating diseases caused by MCH.

15 Also, a compound of the invention is low in toxicity, and is excellent in oral absorbency and intracerebral transitivity.

Therefore, a melanin-concentrating hormone antagonist (hereafter, also abbreviated as "MCH antagonist") comprising a compound of the invention can be 20 safely administered to mammals (e.g. rats, mice, guinea pigs, rabbits, sheep, horses, swine, cattle, monkeys, humans, etc.) as an agent for preventing or treating diseases caused by MCH.

Here, examples of the diseases caused by MCH include 25 obesity (e.g. malignant mastocytosis, exogenous obesity, hyperinsulinar obesity, hyperplasmic obesity, hypophyseal adiposity, hypoplasmic obesity, hypothyroid obesity, hypothalamic obesity, symptomatic obesity, infantile obesity, upper body obesity, alimentary obesity, 30 hypogonadal obesity, systemic mastocytosis, simple obesity, central obesity, etc.), hyperphagia, emotional disorders, reproductive function disorders, memory disorders, dementia, hormonal disorders.

A compound of the invention is also useful as an agent 35 for preventing or treating lifestyle diseases such as diabetes, diabetic complications (e.g. diabetic

retinopathy, diabetic neuropathy, diabetic nephropathy, etc.), arteriosclerosis, and gonitis.

Further, a compound of the invention is useful as an anorectic agent.

5 A MCH antagonist and a pharmaceutical composition of the invention can be used in combination with an alimentary therapy (e.g., alimentary therapy for diabetes) and exercise.

10 A MCH antagonist and a pharmaceutical composition of the invention can be produced by subjecting compound (I) or compound (I') respectively, as it is, or together with a pharmacologically acceptable carrier, to pharmaceutical manufacturing process in accordance with a per se known means.

15 Here, examples of the pharmacologically acceptable carriers include various organic or inorganic carrier substances which are commonly used as materials for pharmaceutical preparations, such as excipients, lubricants, binders, and disintegrators in solid 20 preparations; solvents, solubilizing agents, suspending agents, isotonizing agents, buffering agents, soothing agents, in liquid preparations. Also, in the pharmaceutical manufacturing process, additives such as 25 antiseptics, antioxidants, coloring agents, sweeteners, absorbents, moistening agents, can be used, if necessary.

Examples of the excipients include lactose, sucrose, D-mannitol, starch, cornstarch, crystalline cellulose, light anhydrous silicic acid.

30 Examples of the lubricants include magnesium stearate, calcium stearate, talc, colloidal silica.

Examples of the binders include crystalline cellulose, sucrose, D-mannitol, dextrin, hydroxypropylcellulose, hydroxypropylmethylcellulose, polyvinylpyrrolidone, starch, saccharose, gelatin, 35 methylcellulose, carboxymethylcellulose sodium.

Examples of the disintegrators include starch,

carboxymethylcellulose, carboxymethylcellulose calcium, crosscarmellose sodium, carboxymethylstarch sodium, low-substituted hydroxypropylcellulose (L-HPC).

Examples of the solvents include distilled water for
5 injection, alcohol, propylene glycol, macrogol, sesame oil, corn oil.

Examples of the solubilizing agents include polyethylene glycol, propylene glycol, D-mannitol, benzyl benzoate, ethanol, trisaminomethane, cholesterol,
10 triethanolamine, sodium carbonate, sodium citrate.

Examples of the suspending agents include surfactants such as stearyltriethanolamine, sodium lauryl sulfate, lauryl amino propionic acid, lecithin, benzalkonium chloride, benzethonium chloride, glyceryl monostearate; or
15 hydrophilic polymers such as polyvinyl alcohol, polyvinylpyrrolidone, carboxymethylcellulose sodium, methylcellulose, hydroxymethylcellulose, hydroxyethylcellulose, hydroxypropylcellulose.

Examples of the isotonizing agents include glucose, D-sorbitol, sodium chloride, glycerin, D-mannitol.
20

Examples of the buffering agents include buffer solutions of phosphate, acetate, carbonate and citrate.

Examples of the soothing agents include benzyl alcohol.
25

Examples of the antiseptics include paraoxybenzoates, chlorobutanol, benzyl alcohol, phenethylalcohol, dehydroacetic acid, and sorbic acid.

Examples of the antioxidants include sulfite, ascorbic acid.
30

A MCH antagonist and a pharmaceutical composition of the invention can be safely administered orally or parenterally (e.g. by local, rectal and intravenous administration) in various dosage forms, for instance, as oral drugs such as tablets (including sugar-coated tablets and film-coated tablets), powders, granules, capsules (including soft capsules), solutions; and parenteral

preparations such as injections (e.g. subcutaneous injections, intravenous injections, intramuscular injections, intraperitoneal injections, etc.), external preparations (e.g. nasal preparations, percutaneous preparations, ointments, etc.), suppositories (e.g. rectal suppositories, vaginal suppositories, etc.), sustained-release preparations (e.g. sustained-release microcapsules, etc.), pellets, drip infusions, etc.

The content of compound (I) in a MCH antagonist of the invention and the content of compound (I') in a pharmaceutical composition of the invention are, for instance, about 0.1 to 100 weight percent of the MCH antagonist or whole pharmaceutical composition, respectively.

The dose of a MCH antagonist and a pharmaceutical composition of the invention can be appropriately selected depending on the subject of administration, route of administration, disease, etc.

For instance, the dose per day when a MCH antagonist or a pharmaceutical composition of the invention is orally administered to an adult obesity patient (body weight: about 60 kg), is about 0.1 to about 500 mg, preferably about 1 to about 100 mg, more preferably about 5 to about 100 mg, in terms of compound (I) or compound (I'), each of which is an active ingredient. These amounts can be divided into one to several doses per day for administration.

The MCH antagonist and pharmaceutical composition of the invention can be used in combination with other concomitant drugs which do not interfere with the MCH antagonist and pharmaceutical composition of the invention, for the purpose of "strengthening of therapeutic effect against obesity", "reduction of dose of MCH antagonist", etc. Examples of the concomitant drugs include a "agents for treating diabetes", "agents for treating diabetic complications", "agents for treating obesity other than MCH antagonists", "agents for treating

hypertension", "agents for treating hyperlipidemia (agents for treating arteriosclerosis)", "agents for treating arthritis", "antianxiety agents", "antidepressant". Two or more kinds of these concomitant drugs can be combined
5 in an appropriate ratio for use.

Examples of the above "agents for treating diabetes" include insulin sensitizers, insulin secretion enhancers, biguanides, insulins, α -glucosidase inhibitors, β 3 adrenaline receptor agonists.

10 Examples of the insulin sensitizers include pioglitazone or its salt (preferably hydrochloride), troglitazone, rosiglitazone or its salt (preferably maleate), JTT-501, GI-262570, MCC-555, YM-440, DRF-2593, BM-13-1258, KRP-297, R-119702.

15 Examples of the insulin secretion enhancers include sulfonylureas. Concrete examples of the sulfonylureas include tolbutamide, chlorpropamide, trazamide, acetohexamide, glycopyramide and its ammonium salt, glibenclamide, gliclazide, glimepiride.

20 Other than the above, examples of insulin secretion enhancers include repaglinide, nateglinide, mitiglinide (KAD-1229), JTT-608.

Examples of biguanides include metformin, buformin, phenformin.

25 Examples of insulins include animal insulins extracted from bovine or porcine pancreas; semi-synthetic human insulin which is enzymatically synthesized from insulin extracted from porcine pancreas; human insulin synthesized by genetic engineering, using Escherichia Coli and yeast. As insulin, also employed are insulin-zinc containing 0.45 to 0.9 (w/w)% of zinc; protamine-insulin-zinc produced from zinc chloride, protamine sulfate and insulin. In addition, insulin can be an insulin fragment or derivative (e.g. INS-1, etc.).

30 Insulin can also include various types such as ultra immediate action type, immediate action type, two-phase

type, intermediate type, prolonged action type, etc., and these can be selected depending on the pathological conditions of patients.

Examples of α -glucosidase inhibitors include
5 acarbose, voglibose, miglitol, emiglitate.

Examples of β 3 adrenaline receptor agonists include AJ-9677, BMS-196085, SB-226552, AZ40140.

Other than the above, examples of the "agents for treating diabetes" include ergoset, pramlintide, leptin,
10 BAY-27-9955.

Examples of the above "agents for treating diabetic complications" include aldose reductase inhibitors, glycation inhibitors, protein kinase C inhibitors.

Examples of aldose reductase inhibitors include
15 torulestat; eparlestat; imirestat; zenarestat; SNK-860; zopolrestat; ARI-509; AS-3201.

Examples of glycation inhibitors include pimagedine.

Examples of protein kinase C inhibitors include NGF, LY-333531.

Other than the above, examples of "agents for treating diabetic complications" include alprostadil, thiapride hydrochloride, cilostazol, mexiletine hydrochloride, ethyl eicosapentate, memantine, pimagedline (ALT-711).

Examples of the above "agents for treating obesity other than MCH antagonists" include lipase inhibitors and anorectics.

Examples of lipase inhibitors include orlistat.

Examples of anorectics include mazindol, dextroamphetamine, fluoxetine, sibutramine, baiamine,
30 (S)-sibutramine, SR-141716, NGD-95-1.

Other than the above, examples of "agents for treating obesity other than MCH antagonists" include lipstatin.

Examples of the above "agents for treating hypertension" include angiotensin converting enzyme
35 inhibitors, calcium antagonists, potassium channel openers, angiotensin II antagonists.

Examples of angiotensin converting enzyme inhibitors include captopril, enarapril, alacepril, delapril (hydrochloride), lisinopril, imidapril, benazepril, cilazapril, temocapril, trandolapril, manidipine 5 (hydrochloride).

Examples of calcium antagonists include nifedipine, amlodipine, efonidipine, nicardipine.

Examples of potassium channel openers include levcromakalim, L-27152, AL0671, NIP-121.

10 Examples of angiotensin II antagonists include losartan, candesartan cilexetil, valsartan, irbesartan, CS-866, E4177.

15 Examples of the above "agents for treating hyperlipidemia (agents for treating arteriosclerosis)" include HMG-CoA reductase inhibitors, fibrate compounds.

Examples of HMG-CoA reductase inhibitors include pravastatin, simvastatin, lovastatin, atorvastatin, fluvastatin, lipantil, cerivastatin, itavastatin, ZD-4522, or their salts (e.g. sodium salts, etc.).

20 Examples of fibrate compounds include bezafibrate, clinofibrate, clofibrate, simfibrate.

Examples of the above "agents for treating arthritis" include ibuprofen.

25 Examples of the above "antianxiety agents" include chlordiazepoxide, diazepam, oxazepam, medazepam, cloxazolam, bromazepam, lorazepam, alprazolam, fludiazepam.

30 Examples of the above "antidepressants" include fluoxetine, fluvoxamine, imipramine, paroxetine, sertraline.

The timing of administration of the above concomitant drugs is not limited. The MCH antagonist or pharmaceutical composition and the concomitant drugs can be administrated to the subject simultaneously or at staggered times.

35 The dosages of the concomitant drugs can be determined in accordance with clinically used dosages, and can be

appropriately selected according to the subject of administration, route of administration, diseases and combinations of drugs, etc.

The administration forms for the concomitant drugs are
5 not particularly limited as long as a MCH antagonist or a pharmaceutical composition are used in combination with a concomitant drugs at the time of administration. Examples of such administration forms includes 1) administration of a single preparation obtained by simultaneous preparation
10 of MCH antagonist or pharmaceutical composition together with concomitant drugs, 2) simultaneous administration of two kinds of preparations obtained by separate preparation of MCH antagonist or pharmaceutical composition, and concomitant drugs, through the same route of
15 administration, 3) staggered administration of two kinds of preparations obtained by separate preparation of MCH antagonist or pharmaceutical composition, and concomitant drugs, through the same route of administration, 4)
simultaneous administration of two kinds of preparations
20 obtained by separate preparation of MCH antagonist or pharmaceutical composition, and concomitant drugs, through different routes of administration, 5) staggered administration of two kinds of preparations obtained by separate preparation of MCH antagonist or pharmaceutical composition, and concomitant drugs, through different
25 routes of administration (for instance, administration of MCH antagonist or pharmaceutical composition; and concomitant drugs in this order; or administration in reverse order).

30 The ratio of combination of MCH antagonist or pharmaceutical composition with concomitant drugs can be appropriately selected in accordance with the subject of administration, route of administration and diseases, etc.

35 This invention further relates to "a pharmaceutical comprising a melanin-concentrating hormone antagonist in

combination with at least one species selected from the group consisting of an agent for treating diabetes, an agent for treating hypertension and an agent for treating arteriosclerosis".

5 Here, the "melanin-concentrating hormone antagonist" is not especially limited as long as it is a compound having a melanin-concentrating hormone antagonistic action, and may be either of a peptide compound or a non-peptide compound.

10 As "an agent for treating diabetes", "an agent for treating hypertension" and "an agent for treating arteriosclerosis", those exemplified as the above concomitant drugs can be mentioned.

15 These drugs can be used in the same manner as in the above "combination of MCH antagonist of the invention with concomitant drugs".

20 The pharmaceutical provides excellent effects such as "strengthening of therapeutic effect against obesity", "reduction of dose of MCH antagonist", etc. as compared to single use of each drug.

BEST MODE FOR CARRYING OUT THE INVENTION

25 This invention will be explained further in detail by the following Reference Examples, Examples, Preparation Examples, and Experimental Examples. However, these do not limit this invention, and they can be changed within the scope that does not deviate from the scope of this invention.

30 In the following Reference Examples and Examples, "room temperature" means 0 to 30°C. Anhydrous magnesium sulfate or anhydrous sodium sulfate was used to dry the organic layer. "%" means percent by weight, unless otherwise specified.

35 Infrared absorption spectra were determined by the diffuse reflectance method, using fourier transform type infrared spectrophotometer.

FABMS (pos) is mass spectrum determined by the (+) method, in Fast Atom Bombardment Mass Spectrometry.

Other symbols used in the description have the
5 following meanings.

| | | | |
|----|---------------------|---|---|
| | s | : | singlet |
| | d | : | doublet |
| | t | : | triplet |
| | q | : | quartet |
| 10 | m | : | multiplet |
| | br | : | broad |
| | J | : | coupling constant |
| | Hz | : | Hertz |
| | CDCl ₃ | : | heavy chloroform |
| 15 | DMSO-d ₆ | : | heavy dimethylsulfoxide |
| | THF | : | tetrahydrofuran |
| | DMF | : | N,N-dimethylformamide |
| | DMSO | : | dimethylsulfoxide |
| | WSCD | : | 1-ethyl-3-(3-dimethylaminopropyl) 20 carbodimide |
| 20 | WSC | : | 1-ethyl-3-(3-dimethylaminopropyl) carbodimide hydrochloride |
| | ¹ H-NMR | : | proton nuclear resonance (Free substances were usually measured in |
| 25 | | | CDCl ₃ ,.) |
| | IR | : | infrared absorption spectrum |
| | Me | : | methyl |
| | Et | : | ethyl |
| | HOBT | : | 1-hydroxy-1H-benzotriazole |
| 30 | IPE | : | diisopropyl ether |
| | DMAP | : | 4-dimethylaminopyridine |

In this specification and drawings, when bases and
35 amino acids are shown by codes, these codes are based on
those by the IUPAC-IUB Commission on Biochemical

Nomenclature or common codes in the concerned fields.

Examples of these codes are shown below. Also, where some optical isomers of amino acids can exist, the L form is shown unless otherwise specified.

| | | | |
|----|------|---|-------------------------------------|
| 5 | DNA | : | deoxyribonucleic acid |
| | cDNA | : | complementary deoxyribonucleic acid |
| | A | : | adenine |
| | T | : | thymine |
| | G | : | guanine |
| 10 | C | : | cytosine |
| | RNA | : | ribonucleic acid |
| | mRNA | : | messenger ribonucleic acid |
| | dATP | : | deoxyadenosine triphosphate |
| | dTTP | : | deoxythymidine triphosphate |
| 15 | dGTP | : | deoxyguanosine triphosphate |
| | dCTP | : | deoxycytidine triphosphate |
| | ATP | : | adenosine triphosphate |
| | EDTA | : | ethylenediamine tetraacetic acid |
| | SDS | : | sodium dodecyl sulfate |
| 20 | EIA | : | enzyme immunoassay |
| | Gly | : | glycine |
| | Ala | : | alanine |
| | Val | : | valine |
| | Leu | : | leucine |
| 25 | Ile | : | isoleucine |
| | Ser | : | serine |
| | Thr | : | threonine |
| | Cys | : | cysteine |
| | Met | : | methionine |
| 30 | Glu | : | glutamic acid |
| | Asp | : | aspartic acid |
| | Lys | : | lysine |
| | Arg | : | arginine |
| | His | : | histidine |
| 35 | Phe | : | phenylalanine |
| | Tyr | : | tyrosine |

| | | | |
|----|-----|---|-------------------------------------|
| | Tro | : | tryptophan |
| | Pro | : | proline |
| | Asn | : | asparagine |
| | Gln | : | glutamine |
| 5 | pGl | : | pyroglutamine |
| | Me | : | methyl group |
| | Et | : | ethyl group |
| | Bu | : | butyl group |
| | Ph | : | phenyl group |
| 10 | TC | : | thiazolidine-4(R)-carboxamide group |

Substituents, protecting groups and reagents frequently used in this specification, are shown by the following symbols.

| | | | |
|----|---------------------|---|---|
| 15 | Tos | : | p-toluenesulfonyl |
| | CHO | : | formyl |
| | Bzl | : | benzyl |
| | Cl ₂ Bzl | : | 2,6-dichlorobenzyl |
| | Bom | : | benzyloxymethyl |
| 20 | Z | : | benzyloxycarbonyl |
| | Cl-Z | : | 2-chlorobenzyloxycarbonyl |
| | Br-Z | : | 2-bromobenzyloxycarbonyl |
| | Boc | : | t-butoxycarbonyl |
| | DNP | : | dinitrophenol |
| 25 | Trt | : | trityl |
| | Bum | : | t-butoxymethyl |
| | Fmoc | : | N-9-fluorenylmethoxycarbonyl |
| | HOBt | : | 1-hydroxybenztriazole |
| | HOOBt | : | 3,4-dihydro-3-hydroxy-4-oxo-1,2,3-benzotriazine |
| 30 | HONB | : | 1-hydroxy-5-norbornene-2,3-dicarbodiimide |
| | DCC | : | N,N'-dicyclohexylcarbodiimide |

35 SEQ ID NO in the SEQUENCE LISTING in the specification of the present application shows the following sequences.

[SEQ ID NO : 1] shows a synthetic DNA used for screening of cDNA coding rat SLC-1.

5 [SEQ ID NO : 2] shows a synthetic DNA used for screening of cDNA coding rat SLC-1.

[SEQ ID NO : 3] shows an entire amino acid sequence of rat SLC-1.

10 [SEQ ID NO : 4] shows an entire base sequence of rat SLC-1 cDNA wherein Sal I recognition sequence was added to the 5' side, and Spe I recognition sequence was added to the 3' side.

[SEQ ID NO : 5] shows riboprobe used to determine the quantity of SLC-1 mRNA expressed in each clone of rat SLC-1 expression CHO cells.

15 [SEQ ID NO : 6] shows a synthetic DNA used to obtain cDNA for coding of human SLC-1.

[SEQ ID NO : 7] shows a primer used to make double-strand cDNA for coding human SLC-1.

[SEQ ID NO : 8] shows an entire base sequence of cDNA for coding human SLC-1.

20 [SEQ ID NO : 9] shows an entire amino acid sequence of human SLC-1.

[SEQ ID NO : 10] shows a synthetic DNA used for screening of cDNA for coding human SLC-1(S).

25 [SEQ ID NO : 11] shows a synthetic DNA used for screening of cDNA for coding human SLC-1(S).

[SEQ ID NO : 12] shows a synthetic DNA used for screening of cDNA for coding human SLC-1(L).

30 [SEQ ID NO : 13] shows a synthetic DNA used for screening of cDNA for coding human SLC-1(L).

[SEQ ID NO : 14] shows an entire base sequence of human SLC-1(S) cDNA wherein Sal I recognition sequence was added to the 5' side, and Spe I recognition sequence was added to the 3' side.

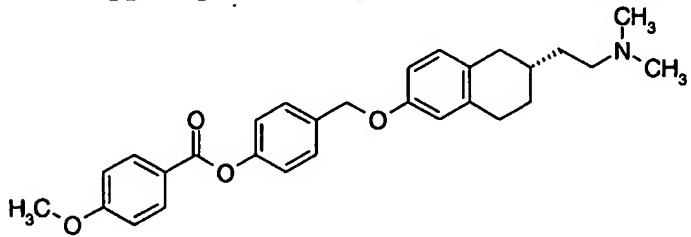
35 [SEQ ID NO : 15] shows an entire base sequence of human SLC-1(L) cDNA wherein Sal I recognition sequence was added

to the 5' side, and Spe I recognition sequence was added to the 3' side.

[SEQ ID NO : 16] shows riboprobe used to determine the quantity of SLC-1mRNA expressed in each clone of human 5 SLC-1(S) expression CHO cells and SLC-1(L) expression CHO cells.

Transformant Escherichia coli DH10B/phSLC1L8 transformed by plasmid containing DNA which codes the base 10 sequence shown by SEQ ID NO : 9, obtained in Reference Example 1 - 6, is on deposit with National Institute of Bioscience and Human-Technology (NIBH), Agency of Industrial Science and Technology, Ministry of International Trade and Industry, as deposit number FERM 15 BP-6632 from February 1, 1999; and with the Institute for Fermentation, Osaka, Japan (IFO), as deposit number IFO 16254 from January 21, 1999.

Reference Example 1
20 2-(R)-[2-(N,N-Dimethylamino)ethyl]-6-[(4-methoxyphenyl)carbonyloxy]benzyloxytetralin



Diethyl azodicarboxylate (40% toluene solution, 0.95 g) was added dropwise to THF solution (6 ml) of 2-(R)- 25 [2-(N,N-dimethylamino)ethyl]-6-hydroxytetralin (300 mg), 4-(hydroxymethyl)phenyl 4-methoxybenzoate (530 mg), and triphenylphosphine (430 mg) under ice-cooling. After stirring for 2 hours at room temperature, the reaction mixture was concentrated. The residue was purified using 30 almina column chromatography (development solvent: hexane ~ hexane : ethyl acetate = 10:1), and the titled compound

(320 mg) was obtained after recrystallization (ethyl acetate-hexane).

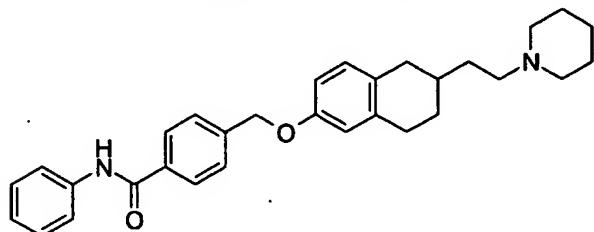
Melting point: 111 - 114°C

$[\alpha]_D^{20} = +44.4^\circ$ (c = 0.502, methanol)

5

Reference Example 2

N-Phenyl-4-[(2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzamide



10 Triethylamine (0.11 ml) was added to THF suspension (3 ml) of 4-[(2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzoate (300 mg). Further, THF solution (0.5 ml) of trimethylacetyl chloride (92 mg) was added dropwise under ice-cooling, which was stirred for 30 minutes. The temperature of the reaction mixture was raised to room temperature, which was stirred for 1 hour. THF solution (0.5 ml) of aniline (85 mg) was added dropwise to the reaction mixture under ice-cooling, which was stirred for 1 hour. After the reaction mixture was stirred for 24 hours at room temperature, saturated sodium bicarbonate solution was added, and extraction was conducted using a mixed solution of ethyl acetate and THF. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized from THF-methanol-IPE to give the titled compound (150 mg).

15 Melting point: 183 - 185°C

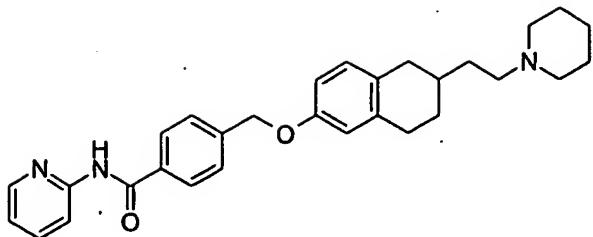
20

25

Reference Example 3

30 4-[(2-(2-Piperidinoethyl)-6-tetralinyl]oxymethyl]-N-(2-pyridinyl)benzamide

90

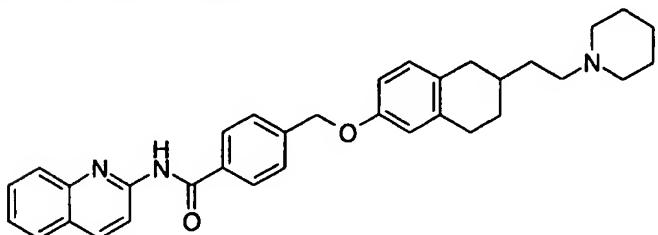


Triethylamine (0.11 ml) was added to THF suspension (6 ml) of 4-[(2-(2-piperidinoethyl)-6-tetralinyl)oxymethyl]benzoate (300 mg). Trimethylacetyl chloride (0.095 ml) was added dropwise to the obtained suspension under ice-cooling, which was stirred for 30 minutes. The temperature of the reaction mixture was raised to room temperature, which was stirred for 1 hour.

10 THF solution (1.0 ml) of 2-aminopyridine (110 mg) was added dropwise to the reaction mixture under ice-cooling, which was stirred for 1 hour. Then the reaction mixture was stirred at room temperature for 6 hours, and at 60°C for 12 hours, which was refluxed with heating for 6 hours.

Saturated sodium bicarbonate solution was added to the 15 reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent: THF), and recrystallized (ethyl acetate-IPE) to give the titled 20 compound (30 mg).

Reference Example 4
25 4-[(2-(2-Piperidinoethyl)-6-tetralinyl)oxymethyl]-N-(2-quinolinyl)benzamide



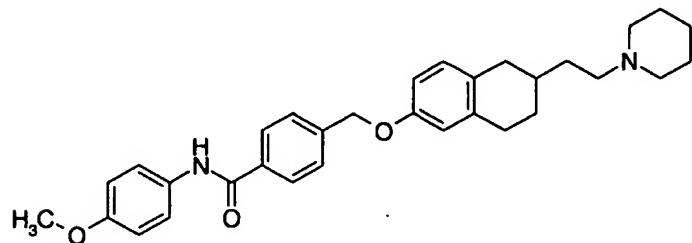
Triethylamine (0.22 ml) was added to THF suspension (6 ml) of 4-[(2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzoate (300 mg). Further, trimethylacetyl chloride (0.095 ml) was added dropwise to 5 under ice-cooling, which was stirred for 30 minutes. The temperature of the reaction mixture was raised to room temperature, which was stirred for 1 hour. THF solution (1.0 ml) of 2-aminoquinoline (170 mg) was added dropwise to the reaction mixture under ice-cooling, which was 10 stirred at room temperature for 12 hours. Saturated sodium bicarbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The 15 residue was purified using alumina column chromatography (development solvent: THF), and recrystallized (ethyl acetate-diisopropyl ether) to give the titled compound (45 mg).

Melting point: 135 - 138°C

20

Reference Example 5

N-(4-Methoxyphenyl)-4-[(2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzamide



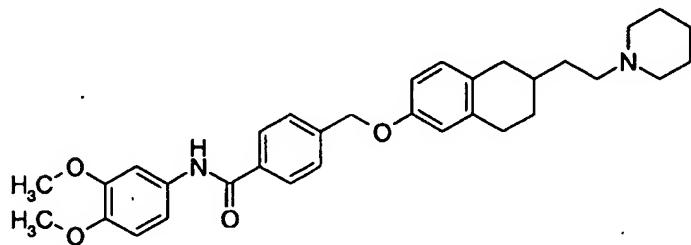
25 WSCD (0.11 ml) was added to DMF solution (2 ml) of 4-[(2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzoate (170 mg), 4-methoxyaniline (53 mg), HOBt (70 mg) and DMAP (60 mg) at room temperature, which was stirred for 12 hours. 10% aqueous potassium 30 carbonate solution and water was added to the reaction mixture, and extraction was conducted using a mixed

solution of THF and ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent: THF), and recrystallized (THF-IPE) to give the titled compound (140 mg).

Melting point: 193 - 196°C

Reference Example 6

10 N-(3,4-Dimethoxyphenyl)-4-[[2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzamide

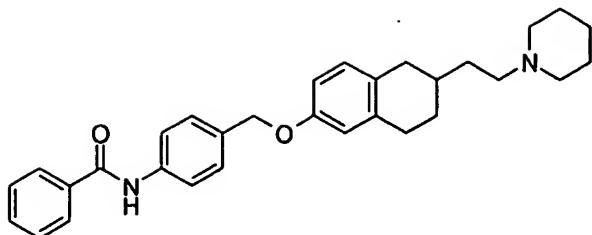


WSCD (free form, 0.2 ml) was added to DMF solution (3 ml) of 4-[[2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzoate (300 mg), 3,4-dimethoxyaniline (120 mg), HOBT (120 mg) and DMAP (100 mg) at room temperature, which was stirred for 12 hours. 10% aqueous potassium carbonate solution was added to the reaction mixture, and the resulting crystals were collected by filtration. The crystals were washed with water, then dried. The crystals were purified using alumina column chromatography (development solvent: THF), and recrystallized (THF-IPE) to give the titled compound (330 mg).

25 Melting point: 178 - 180°C

Reference Example 7

6-[4-(Benzoylamino)benzyloxy]-2-(2-piperidinoethyl)tetralin



Sodium hydride (60% oily, 85 mg) was added to DMF solution of 6-hydroxy-2-(2-piperidinoethyl)tetralin (500 mg) at room temperature, which was stirred for 1 hour.

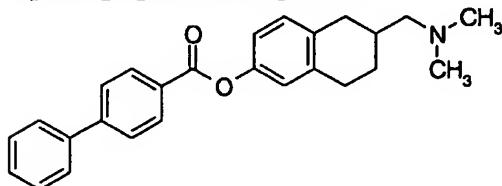
5 N-[4-(bromomethyl)phenyl]benzamide (670 mg) was added to the reaction mixture at room temperature, which was stirred for 1 hour. Water was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent : THF), and recrystallized (ethyl acetate) to give the titled compound (200 mg).

Melting point: 176 - 179°C

15

Reference Example 8

2-[(N,N-Dimethylamino)methyl]-6-tetralinyl 4-biphenylylcarboxylate



20 4-Biphenylylcarboxylic acid (580 mg) and WSC (560 mg) were added to pyridine solution (6 ml) of 2-[(N,N-dimethylamino)methyl]-6-hydroxytetralin (300 mg), which was stirred at room temperature for 36 hours. Saturated sodium bicarbonate solution and water were added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then

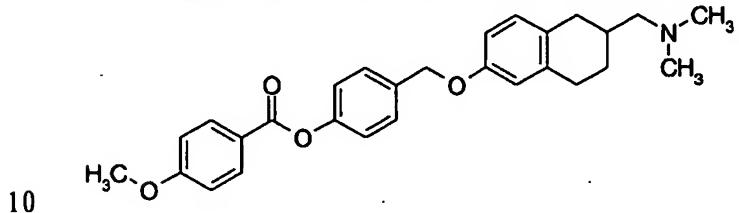
25

concentrated. The residue was purified using alumina column chromatography (development solvent; hexane - hexane:ethyl acetate = 10:1), and recrystallized (hexane) to give the titled compound (300 mg).

5 Melting point: 85 - 86°C

Reference Example 9

2-[(N,N-Dimethylamino)methyl]-6-[4-[(4-methoxyphenyl)carbonyloxy]benzyloxy]tetralin



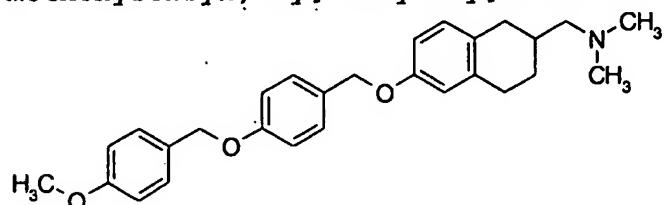
10

Diethyl azodicarboxylate (40% toluene solution, 950 mg) was added dropwise to THF solution (3 ml) of 2-[(N,N-dimethylamino)methyl]-6-hydroxytetralin (150 mg), 4-(hydroxymethyl)phenyl 4-methoxybenzoate (570 mg) and triphenylphosphine (574 mg) at room temperature, which was stirred for 3 hours. The reaction mixture was concentrated, and the residue was purified using alumina column chromatography (development solvent; hexane - hexane:ethyl acetate = 6:1), and recrystallized (ethyl acetate-hexane) to give the titled compound (175 mg).

20 Melting point: 119 - 121°C

Reference Example 10

2-[(N,N-Dimethylamino)methyl]-6-[4-[(4-methoxybenzyl)oxy]benzyloxy]tetralin



25 Diethyl azodicarboxylate (40% toluene solution, 1.91 g) was added dropwise to THF solution (6 ml) of 2-

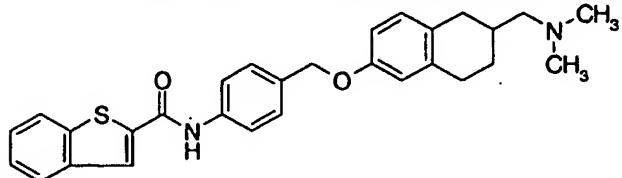
[(N,N-dimethylamino)methyl]-6-hydroxytetralin (300 mg), 4-[(4-methoxybenzyl)oxy]benzylalcohol (1.07 g) and triphenylphosphine (1.15g) at room temperature, which was stirred for 12 hours. The reaction mixture was concentrated, and the residue was purified using alumina column chromatography (development solvent; hexane ~ hexane:ethyl acetate = 10:1), and recrystallized (ethyl acetate-hexane) to give the titled compound (260 mg).

Melting point: 106 - 111°C

10

Reference Example 11

6-[4-[(1-Benzothiophen-2-yl)carbonylamino]benzyloxy]-2-[(N,N-dimethylamino)methyl]tetralin

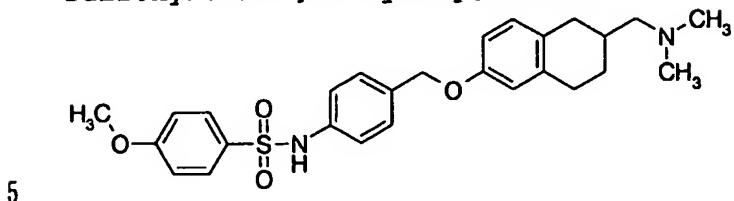


15 One drop of DMF was added to THF solution (4 ml) of 1-benzothiophene-2-carboxylic acid (230 mg), and oxalyl chloride (0.23 ml) was further added under ice-cooling, which was stirred for 30 minutes at room temperature. The reaction mixture was concentrated, which was dissolved in 20 THF (1 ml). The obtained solution was added dropwise to pyridine solution (6 ml) of 6-(4-aminobenzyloxy)-2-[(N,N-dimethylamino)methyl]tetralin (300 mg), which was stirred for 15 minutes. After stirring at room temperature for another 15 minutes, 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development 25 solvent; ethyl acetate), and recrystallized (THF-IPE) to give the titled compound (250 mg).

Melting point: 165 - 169°C

Reference Example 12

2-[(N,N-Dimethylamino)methyl]-6-[4-[(4-methoxyphenyl)sulfonylamino]benzyloxy]tetralin

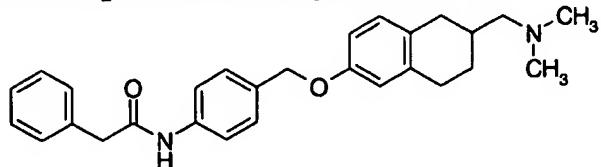


THF solution (1 ml) of 4-methoxybenzenesulfonyl chloride (270 mg) was added dropwise to pyridine solution (6 ml) of 6-[(4-aminobenzyl)oxy]-2-[(N,N-dimethylamino)methyl]tetralin (300 mg) under ice-cooling, which was stirred for 15 minutes. After stirring at room temperature for further 15 minutes, 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent: ethyl acetate), and recrystallized (ethyl acetate-IPE) to give the titled compound (260 mg). Melting point: 137 - 140°C.

20

Reference Example 13

6-[(4-Benzylcarbonylamino)benzyloxy]-2-[(N,N-dimethylamino)methyl]tetralin



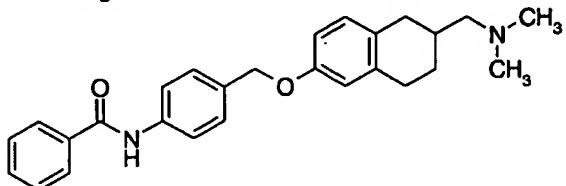
25 THF solution (1 ml) of phenylacetyl chloride (200 mg) was added dropwise to pyridine solution (6 ml) of 6-[(4-aminobenzyl)oxy]-2-[(N,N-dimethylamino)methyl]tetralin (300 mg) under ice-cooling, which was stirred for 15 minutes. After stirring at room temperature for further 15 minutes, saturated sodium

30

bicarbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent; hexane - hexane:ethyl acetate = 2:1), and recrystallized to give the titled compound (175 mg).
Melting point: 130 - 135°C

10 Reference Example 14

6-[4-(Benzoylamino)benzyloxy]-2-[(N,N-dimethylamino)methyl] tetralin

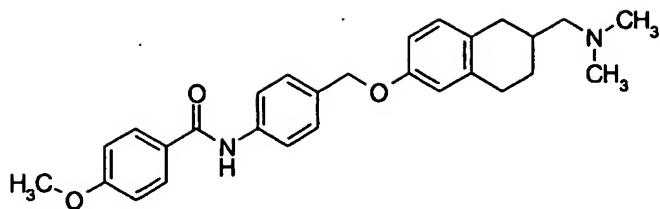


Benzoyl chloride (0.14 ml) was added dropwise to pyridine solution (6 ml) of 6-[(4-aminobenzyl)oxy]-2-[(N,N-dimethylamino)methyl]tetralin (300 mg) under ice-cooling, which was stirred at room temperature for 30 minutes. 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent; ethyl acetate), and recrystallized (THF-IPE) to give the titled compound (240 mg).

Melting point: 128 - 133°C

Reference Example 15

2-[(N,N-Dimethylamino)methyl]-6-[4-[(4-methoxybenzoyl)amino]benzyloxy]tetralin



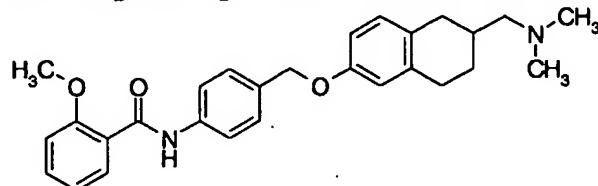
p-Anisoyl chloride (0.20 ml) was added dropwise to pyridine solution (6 ml) of 6-[(4-aminobenzyl)oxy]-2-[(N,N-dimethylamino)methyl]tetralin (300 mg) under ice-cooling, which was stirred at room temperature for 30 minutes. 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using alumina column chromatography (development solvent: ethyl acetate), and recrystallized (THF-IPE) to give the titled compound (300 mg).

Melting point: 155 - 159°C

15

Reference Example 16

2-[(N,N-Dimethylamino)methyl]-6-[4-[(2-methoxybenzoyl)amino]benzyloxy]tetralin



20 o-Anisoyl chloride (0.15 ml) was added dropwise to pyridine solution (4 ml) of 6-[(4-aminobenzyl)oxy]-2-[(N,N-dimethylamino)methyl]tetralin (200 mg) under ice-cooling, which was stirred at room temperature for 30 minutes. 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified

25

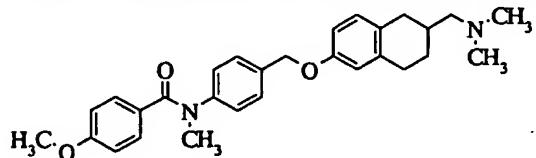
using alumina column chromatography (development solvent; THF), and recrystallized (ethyl acetate-hexane) to give the titled compound (200 mg).

Melting point: 106 - 108°C

5

Reference Example 17

6-[4-[N-(4-Methoxybenzoyl)-N-methylamino]benzyloxy]-2-[(N,N-dimethylamino)methyl]tetralin

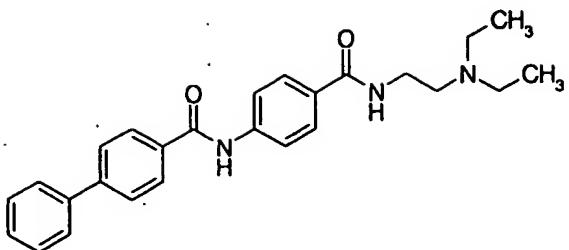


10 Diethyl azodicarboxylate (40% toluene solution, 960 mg) was added dropwise to THF solution (3 ml) of 2-[(N,N-dimethylamino)methyl]-6-hydroxytetralin (150 mg), N-[4-(hydroxymethylphenyl)-4-methoxy-N-methylbenzamide (600 mg) and triphenylphosphine (570 mg) at room
 15 temperature, which was stirred for 12 hours. After the reaction mixture was concentrated, the residue was purified using silica gel column chromatography (development solvent; hexane ~ ethyl acetate ~ ethyl acetate:methanol = 1:2), and then purified using alumina column
 20 chromatography (development solvent; hexane - hexane:ethyl acetate = 2:1) to give the titled compound (185 mg).
¹H-NMR (CDCl₃) δ : 1.20-1.50(1H, m), 1.80-2.46(5H, m), 2.25(6H, s), 2.68-2.86(3H, m), 3.47(3H, s), 3.74(3H, s), 4.95(2H, s), 6.52-6.76(4H, m), 6.84-7.14(3H, m), 7.22-25 7.38(4H, m).

Reference Example 18

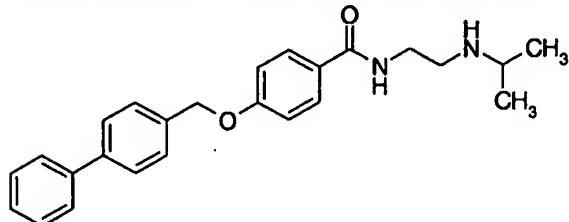
N-[4-[[[2-(Diethylamino)ethyl]amino]carbonyl]phenyl] 4-biphenylylcarboxamide

100



Oxalyl chloride (0.46 ml) and DMF (1 drop) were added to THF solution (15 ml) of 4-biphenylcarboxylic acid (0.879g) under ice-cooling. The reaction mixture was
 5 stirred at room temperature for 30 minutes, and concentrated. The residue was dissolved in THF (10 ml), which was added dropwise to THF (20 ml) suspension of procaineamide hydrochloride (1.078 g) and triethylamine (1.4 ml) at 0°C. After stirring at 0°C for 30 minutes, 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized using
 10 methanol to give the titled compound (1.147 g).
 15 Melting point: 237 - 240°C (decomposition)

Reference Example 19
 4-(4-Biphenylmethoxy)-N-[2-(isopropylamino)ethyl]benzamide



WSC (0.708 g), HOBt (0.521 g), N-isopropyl
 20 ethylenediamine (0.353 g) and triethylamine (1 ml) were added to a mixed solution of 4-(4-biphenylmethoxy)
 25 benzoate (1.007 g) in THF (30 ml) and acetonitrile (30 ml). After stirring at room temperature for 18 hours, water was added to the reaction mixture, and extraction was conducted

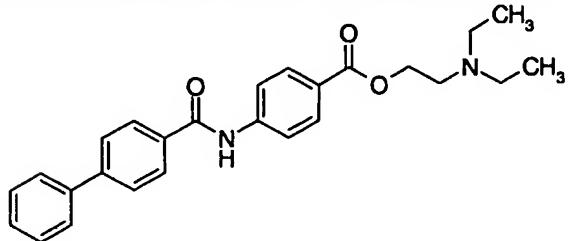
using ethyl acetate. The organic layer was washed with 10% aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried, and then concentrated.

The residue was recrystallized using ethanol to give the
5 titled compound (0.806 g).

Melting point: 150 - 154°C

Reference Example 20

10 2-(N,N-Diethylamino)ethyl 4-(4-
biphenylylcarbonylamino)benzoate



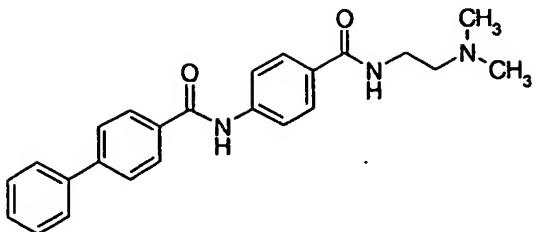
Oxalyl chloride (0.39 ml) and DMF (1 drop) were added to THF solution (15 ml) of 4-biphenylylcarboxylic acid (1.091 g) under ice-cooling, which was stirred at room
15 temperature for 30 minutes, and concentrated. The residue was dissolved in THF (10 ml), which was added dropwise to THF suspension (30 ml) of procaine hydrochloride (1.091 g) and triethylamine (0.67 ml) at 0°C. After stirring at 0°C for 30 minutes, 10% aqueous potassium carbonate was added
20 to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized using ethyl acetate/hexane to give the titled compound (0.728 g).

25 Melting point: 146 - 149°C

Reference Example 21

N-[4-[[[2-(Dimethylamino)ethyl]amino]carbonyl]phenyl]
4-biphenylylcarboxamide

102



WSC (0.248 g), HOBT (0.156 g), N,N-dimethyl ethylenediamine (0.097 g) and triethylamine (0.21 ml) were added to a mixed solution of 4-(4-biphenylcarbonylamino)benzoate (0.323 g) in THF (15 ml) and acetonitrile (15 ml). After stirring at room temperature for 18 hours, water was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with 10% aqueous potassium carbonate and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized using methanol/diethyl ether to give the titled compound (0.100 g).

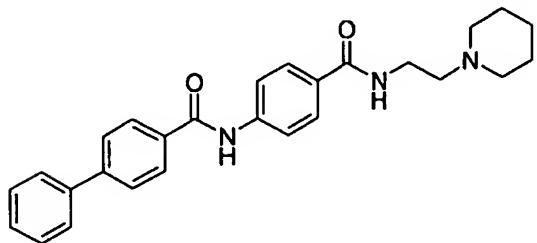
Melting point: 261 - 264°C (decomposition)

15

The compounds described in the following Reference Examples 22 to 25 were produced in the same manner as in Reference Example 21.

20 Reference Example 22

N-[4-[(2-(Piperidinoethyl)amino]carbonyl]phenyl] 4-biphenylcarboxamide

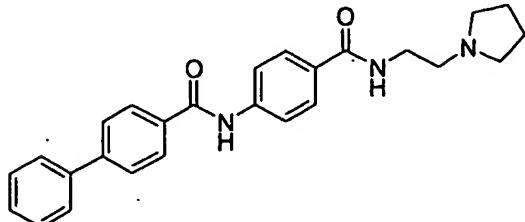


Melting point: 247 - 252°C (decomposition)

25

Reference Example 23

**N-[4-[[2-(1-Pyrrolidinyl)ethyl]amino]carbonyl]phenyl]
4-biphenylylcarboxamide**

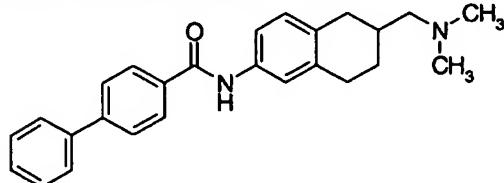


Melting point: 241 - 245°C (decomposition)

5

Reference Example 24

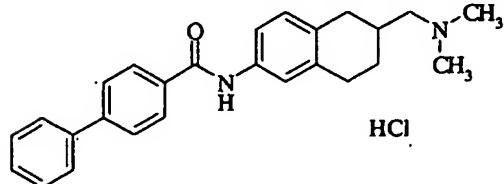
**N-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-4-
biphenylylcarboxamide**



10 Melting point: 164 - 166°C

Reference Example 25

**N-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-4-
biphenylylcarboxamide hydrochloride**



15

Melting point: >250°C

$^1\text{H-NMR}$ δ : 1.24-1.54 (1H, m), 1.84-2.10 (2H, m), 2.20-2.50
(3H, m), 2.26 (6H, s), 2.79-3.01 (3H, m), 7.10 (1H, d,

$J=8\text{Hz}$), 7.28-7.54 (5H, m), 7.60-7.82 (5H, m), 7.94 (2H, d,

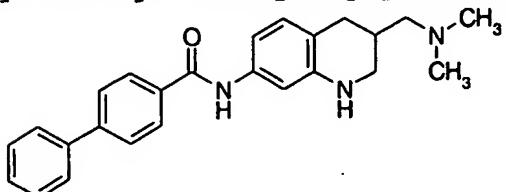
20 $J=8\text{Hz}$).

IR(KBr) 3028, 2910, 2640, 1658, 1538, 1417, 746, 701 cm^{-1}

Reference Example 26

N-[3-[(N,N-Dimethylamino)methyl]-1,2,3,4-tetrahydro-7-

quinolinyl]-4-biphenylylcarboxamide



One drop of DMF was added to THF solution of 4-biphenylylcarboxylic acid (145 mg), and oxalyl chloride (0.1 ml) was added dropwise to the solution under ice-cooling, which was stirred at room temperature for 30 minutes. After the reaction mixture was concentrated, the residue was dissolved in THF (1 ml), which was added dropwise to pyridine solution (1.5 ml) of 7-amino-3-[(N,N-dimethylamino)methyl]-1,2,3,4-tetrahydroquinoline (150 mg) under ice-cooling, and the reaction mixture was stirred for 30 minutes. After the temperature of the reaction mixture was raised to room temperature, 10% aqueous potassium carbonate was added to the reaction mixture, and extraction was conducted using a mixed solution of THF and ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized using THF-IPE to give the titled compound (180 mg).

Melting point: 206 - 211°C

Reference Example 27

25 4-[N-[(Benzyl)carbonyl]-N-methylamino]-N-[3-[(N,N-dimethylamino)methyl]-1,2,3,4-tetrahydro-7-quinolinyl]benzamide



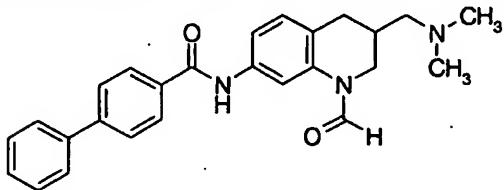
One drop of DMF was added dropwise to THF solution (2

ml) of 4-[N-[(benzyloxy)carbonyl]-N-methylamino]benzoic acid (210 mg), and then oxalyl chloride (0.1 ml) was added dropwise under ice-cooling, which was stirred at room temperature for 30 minutes. After the reaction mixture was 5 concentrated, the residue was dissolved in THF (1 ml), which was added dropwise to pyridine solution (1.5 ml) of 7-amino-3-[(N,N-dimethylamino)methyl]-1,2,3,4-tetrahydroquinoline (150 mg) under ice-cooling. The reaction mixture was then stirred for 30 minutes. After 10 the temperature of the reaction mixture was raised to room temperature, 10% aqueous potassium carbonate solution was added, and extraction was conducted using a mixed solution of THF and ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, 15 dried, and then concentrated. The residue was recrystallized using THF-IPE to give the titled compound (220 mg).

Melting point: 167 - 172°C

20 Reference Example 28

N-[3-[(N,N-Dimethylamino)methyl]-1-formyl-1,2,3,4-tetrahydron-7-quinolinyl]-4-biphenylylcarboxamide



Anhydrous acetic acid (0.1 ml) was added to formic acid 25 (1 ml), which was stirred at 55°C for 2 hours. N-[3-[(N,N-dimethylamino)methyl]-1,2,3,4-tetrahydron-7-quinolinyl]-4-biphenylylcarboxamide (80 mg) was added to the reaction mixture under ice-cooling, which was stirred at room temperature for 72 hours. 10% aqueous potassium carbonate solution was added to the reaction mixture to make the mixture alkaline, and extraction was conducted using ethyl acetate. The organic layer was washed with water and 30

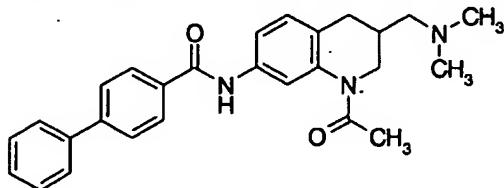
saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized using THF-IPE to give the titled compound (80 mg).

Melting point: 134 - 138°C

5

Reference Example 29

N-[1-Acetyl-3-[(N,N-dimethylamino)methyl]-1,2,3,4-tetrahydro-7-quinolyl]-4-biphenylylcarboxamide

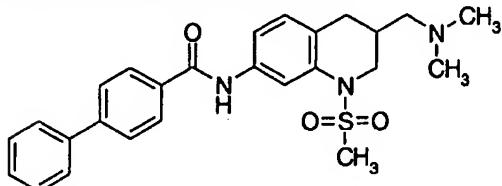


10 Acetyl chloride(0.02 ml) was added to pyridine solution (1 ml) of N-[3-[(N,N-dimethylamino)methyl]-1,2,3,4-tetrahydro-7-quinolyl]-4-biphenylylcarboxamide (80 mg) under ice-cooling, which was stirred for 15 minutes, and then stirred at room temperature
15 for 15 minutes. 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was
20 recrystallized using THF-IPE to give the titled compound (64 mg).

Melting point: 167 - 173°C

Reference Example 30

25 **N-[3-[(N,N-Dimethylamino)methyl]-1-methylsulfonyl-1,2,3,4-tetrahydro-7-quinolyl]-4-biphenylylcarboxamide**



Methanesulfonyl chloride (0.02 ml) was added to

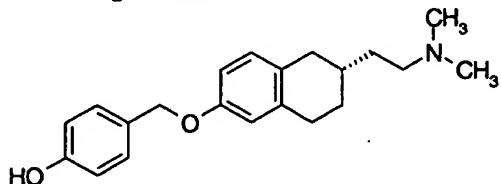
pyridine solution (1 ml) of N-[3-[(N,N-dimethylamino)methyl]-1,2,3,4-tetrahydro-7-quinolinyl]-4-biphenylcarboxamide (80 mg) under ice-cooling, which was stirred at room temperature for 1 hour. Further, 5 methanesulfonyl chloride (0.02 ml) was added to the reaction mixture under ice-cooling, which was stirred at room temperature for 12 hours. 10% aqueous potassium carbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic 10 layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized using THF-IPE to give the titled compound (64 mg).

Melting point: 184 - 188°C

15

Reference Example 31

2-(R)-[2-(N,N-Dimethylamino)ethyl]-6-(4-hydroxyphenyl)methoxytetralin



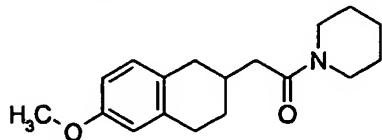
20 THF solution (2 ml) of 2-(R)-[2-(N,N-dimethylamino)ethyl]-6-[4-(4-methoxyphenylcarbonyloxy)phenylmethoxy]tetralin (330 mg) was added dropwise to THF suspension (4 ml) of lithium aluminum hydride (60 mg) under ice-cooling. 1N aqueous sodium hydroxide solution was 25 added to the reaction mixture to make the mixture basic, and the precipitate was removed by celite filtration. After the filtrate was concentrated, the residue was purified using silica gel chromatography (development solvent; ethyl acetate - methanol), and recrystallized (ethyl 30 acetate-hexane) to give the titled compound (70 mg).

Melting point: 132 - 135°C

$[\alpha]_D^{20} = +56.9^\circ$ (c = 0.505, methanol)

Reference Example 32

2-(6-Methoxy-2-tetralinyl)-1-piperidino-1-ethanone

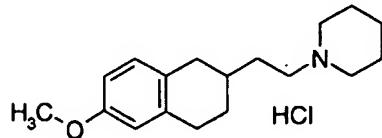


5 2-(6-Methoxy-2-tetralinyl)acetic acid (8.8 g) was dissolved in a mixed solution of THF (150 ml) and acetonitrile (50 ml), then piperidine (5.2 g), WSC (12 g), HOBT (6.0 g) and triethylamine (17 ml) were added to the solution, which was stirred at room temperature for 12 hours. Water was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with 1N hydrochloric acid, water, saturated sodium bicarbonate solution, water, and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using silica gel chromatography (development solvent; ethyl acetate) to give the titled compound (10.3 g). Recrystallization from hexane gave crystals of the following melting points.
Melting point: 59 - 61°C

20

Reference Example 33

6-Methoxy-2-(2-piperidinoethyl)tetralin hydrochloride



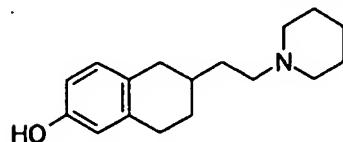
25 THF solution (50 ml) of 2-(6-methoxy-2-tetralinyl)-1-piperidino-1-ethanone (9.80 g) was added dropwise to THF suspension (100 ml) of lithium aluminum hydride (1.94 g) under ice-cooling. The temperature of the reaction mixture was raised to 60°C over 30 minutes, which was stirred for 30 minutes. After the reaction mixture was cooled to room temperature, 1N aqueous sodium hydroxide solution was added to make the reaction mixture basic, and

the precipitate was removed by celite filtration. The filtrate was concentrated and the residue was made into a hydrochloride, which was then recrystallized from ethanol-IPE to give the titled compound (9.80 g).

5 Melting point: 189 - 191°C

Reference Example 34

6-Hydroxy-2-(2-piperidinoethyl)tetralin

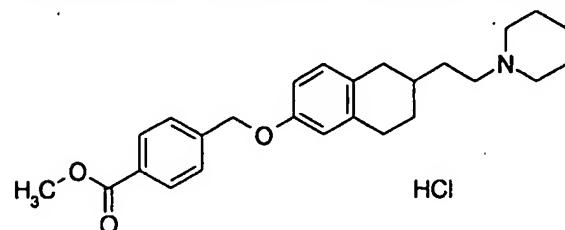


10 **6-Methoxy-2-(2-piperidinoethyl)tetralin**
hydrochloride (9.3 g) was added to 48% hydrobromic acid (50 ml), which was refluxed with heating for 4 hours. After the reaction mixture was concentrated under reduced pressure, saturated sodium bicarbonate solution was added
15 to the residue to make the water layer alkaline, and the water layer was extracted using a mixed solution of THF and ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting crystal was washed with IPE
20 to give the titled compound (5.8 g).

Melting point: 154 - 157°C

Reference Example 35

Methyl 4-[(2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzoate hydrochloride



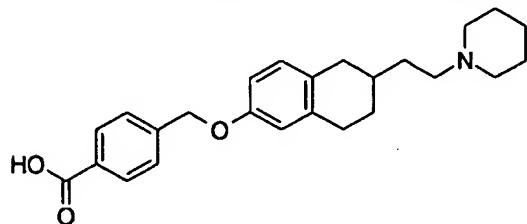
Diethyl azodicarboxylate (40% toluene solution, 5.10 g) was added dropwise to THF solution (15 ml) of 6-hydroxy-2-(2-piperidinoethyl)tetralin (1.50 g), methyl

4-(hydroxymethyl)benzoate (1.44 g), and triphenylphosphine (2.60 g) at room temperature, which was stirred for 12 hours, and then concentrated. The residue was purified using aluminum column chromatography
5 (development solvent; hexane ~ hexane:ethyl acetate = 15:1), which was made into a hydrochloride. The hydrochloride was recrystallized (methanol-IPE) to give the titled compound (1.36 g).

Melting point: 190 - 193°C.
10

Reference Example 36

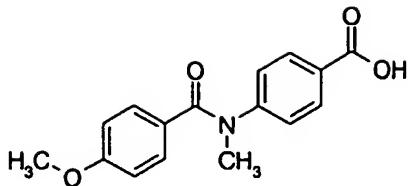
4-[[2-(2-Piperidinoethyl)-6-tetralinyl]oxymethyl]benzoic acid



15 3N Aqueous sodium hydroxide solution (1.8 ml) was added to methanol solution (20 ml) of methyl 4-[[2-(2-piperidinoethyl)-6-tetralinyl]oxymethyl]benzoate hydrochloride (1.06 g), which was refluxed with heating for 6 hours. After the reaction mixture was concentrated,
20 water was added to the reaction mixture. Further, 1N hydrochloric acid was added to make the pH of the mixture about 7. The resulting crystals were filtered to give the titled compound (0.93 g). Recrystallization from ethanol gave crystals of the following melting points.
25 Melting point: 105 - 108°C

Reference Example 37

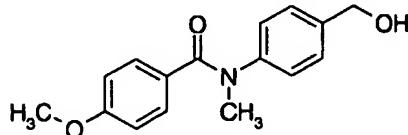
4-[N-(4-Methoxybenzoyl)-N-methylamino]benzoic acid



Aqueous solution (50 ml) of sodium carbonate (23 g) was added to THF solution (50 ml) of 4-(methylamino)benzoic acid (5.0 g), and p-anisoyl chloride (5.6 g) was added dropwise to the solution under ice-cooling, which was stirred for 15 minutes, and then stirred at room temperature for 30 minutes. Concentrated hydrochloric acid was added to the reaction mixture under ice-cooling to make the water layer acidic, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using silica gel column chromatography (development solvent; hexane ~ hexane:ethyl acetate = 1:2), and recrystallized (ethyl acetate-hexane) to give the titled compound (4.8 g).
 Melting point: 157 - 160°C.

Reference Example 38

N-[4-(Hydroxymethyl)phenyl]-4-methoxy-N-methylbenzamide



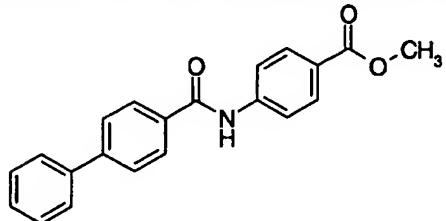
THF solution (1M, 16 ml) of borane was added dropwise to THF solution (10 ml) of 4-[N-(4-methoxybenzoyl)-N-methylamino]benzoic acid (1.14 g) under ice-cooling, which was stirred for 15 minutes, and then stirred at room temperature for 1 hour. After water was added to the reaction mixture, 1N hydrochloric acid was further added, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated sodium bicarbonate, and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was

purified using silica gel chromatography (development solvent; hexane ~ hexane:ethyl acetate = 1:2), and recrystallized (ethyl acetate-hexane) to give the titled compound (770 mg).

5 Melting point: 85 - 90°C.

Reference Example 39

Methyl 4-(4-biphenylylcarbonylamino)benzoate



10 Oxalyl chloride (1.2 ml) and DMF (0.04 ml) were added to THF solution (30 ml) of 4-biphenylylcarboxylic acid (2.184g) under ice-cooling. The reaction mixture was stirred at room temperature for 30 minutes, which was concentrated. The residue was dissolved in THF (15 ml),

15 which was added dropwise to THF solution (30 ml) of methyl 4-aminobenzoate (1.512 g) and triethylamine (2.1 ml) at 0°C. After the reaction mixture was stirred at 0°C for 30 minutes, 10% citric acid solution was added to the reaction mixture, and extraction was conducted using ethyl acetate.

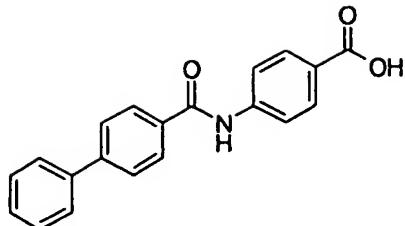
20 The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting crude crystal was washed with diethyl ether to give the titled compound (2.179 g).

Melting point: 247 - 251°C.

25

Reference Example 40

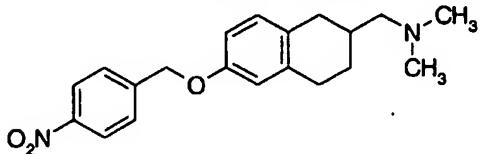
4-(4-Biphenylylcarbonylamino)benzoic acid



1N Aqueous sodium hydroxide solution (8 ml) was added to a mixed solution of methyl 4-(4-biphenylcarbonylamino)benzoate (1.998 g) in THF (60 ml) and methanol (20 ml), which was stirred at room temperature for 18 hours. 1N Hydrochloric acid (10 ml) was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting crude crystals were washed with diethyl ether to give the titled compound (1.760 g).
Melting point: >320°C.
¹H NMR (DMSO-d₆) δ: 7.37-7.57 (3H,m), 7.77 (2H,d), 7.85 (2H,d), 7.95 (4H,s), 8.08 (2H,d), 10.56 (1H,s)

15

Reference Example 41
2-[(N,N-Dimethylamino)methyl]-6-(4-nitrobenzyloxy)tetralin

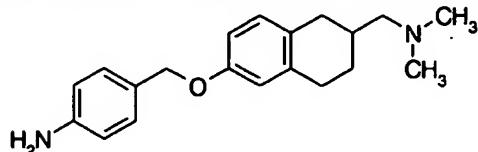


20 Diethyl azodicarboxylate (40% toluene solution, 9.53 g) was added dropwise to THF solution (15 ml) of 2-[(N,N-dimethylamino)methyl]-6-hydroxytetralin (1.5 g), 4-nitrobenzylalcohol (3.35 g), and triphenylphosphine (5.74 g) at room temperature, which was stirred for 24 hours. The reaction mixture was concentrated, and the residue was purified using alumina column chromatography (development solvent; hexane ~ hexane:ethyl acetate = 8:1), and recrystallized (ethyl acetate-hexane) to give the titled compound (1.29 g).

Melting point: 83 - 89°C

Reference Example 42

6-(4-Aminobenzyloxy)-2-[(N,N-dimethylamino)methyl]tetralin



After acetic acid (6 ml) was added to THF solution (12 ml) of 2-[(N,N-dimethylamino)methyl]-6-(4-nitrobenzyloxy)tetralin (1.91 g) under ice-cooling, zinc powder (3.67 g) was further added, which was stirred for 6 hours. The reaction mixture was filtered, and the filtrate was concentrated. 10% aqueous potassium carbonate solution and ethyl acetate were added to the residue, the precipitate was removed by celite filtration, and the filtrate was extracted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was purified using aluminum column chromatography (development solvent; hexane ~ hexane:ethyl acetate = 4:1) to give the titled compound (1.05 g).

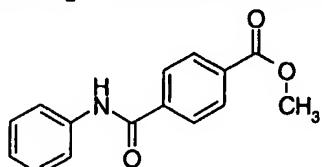
Amorphous powder:

$^1\text{H-NMR}$ (CDCl_3) δ : 1.18-1.50(1H, m), 1.70-2.50(5H, m), 2.24(6H, s), 2.72-2.86(3H, m), 3.68(2H, brs), 4.88(2H, s), 6.58-6.82(4H, m), 6.99(1H, s), 7.14-7.30(2H, m).

25

Reference Example 43

Methyl 4-anilinocarbonylbenzoate



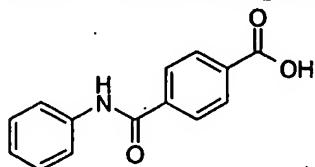
4-Methoxycarbonyl benzoic acid (540 mg), aniline (0.27 ml), WSC (863 mg) and triethylamine (0.84 ml) were

added to THF (20 ml). After the reaction mixture was stirred at room temperature for 20 hours, the reaction mixture was placed in water, and extraction was conducted using ethyl acetate-THF (1:1). The organic layer was 5 washed with water, saturated sodium bicarbonate solution, and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting crude crystals were recrystallized from ethyl acetate-hexane to give the titled compound (659 mg).

10 Melting point: 189 - 190°C

Reference Example 44

4-Anilinocarbonylbenzoic acid

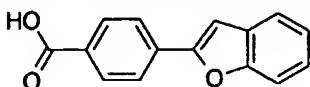


15 8 mol of aqueous sodium hydroxide solution (8 ml) was added to methanol (16 ml) - THF (6 ml) solution of 4-methyl anilinocarbonylbenzoate (511 mg), which was stirred at room temperature for 1 hour. 1 mol of hydrochloric acid was added to the reaction mixture to make the pH of the mixture 20 to 5, extraction was conducted using ethyl acetate-THF (1:1). The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting residue was washed with hexane to give the titled compound (480 mg).

25 Melting point: 305 - 307°C.

Reference Example 45

4-(2-Benzo[b]furanyl)benzoic acid



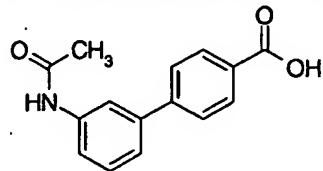
30 Benzofuranyl-2-boric acid (2.1 g), palladium tetratriphenylphosphine (200 mg) and 2M aqueous sodium

carbonate solution were added to toluene (40 ml) - ethanol (10 ml) solution of ethyl 4-bromobenzoate (2.3 g), which was refluxed at 80°C for 5 hours under an argon atmosphere.

The reaction mixture was diluted with water, and
5 extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting residue was purified using silica gel chromatography (development solvent; ethyl acetate:hexane
10 = 1:4), and concentrated, which was dissolved in methanol (10 ml) - THF (10 ml). 8 mol of aqueous sodium hydroxide solution (8 ml) was added to the resulting solution at room temperature, which was stirred for 2 hours. After 1 mol of hydrochloric acid was added to the reaction mixture to
15 make the mixture acidic, extraction was conducted using ethyl acetate-THF (1:1). The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting residue was washed with hexane to give the titled compound (2.272 g).
20 Melting point: 292 - 294°C.

Reference Example 46

3'-Acetamino-4-biphenylcarboxylic acid

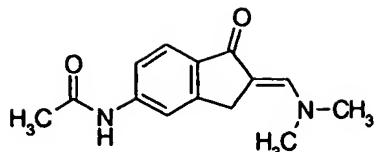


25 The titled compound was produced in the same manner as in Reference Example 45.

Melting point: 300 - 301°C

Reference Example 47

30 N-[2-[(E)-(Dimethylamino)methylidene]-1-oxo-2,3-dihydro-1H-inden-5-yl]acetamide



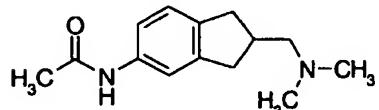
Dimethylformamide dimethylacetal was added to 5-acetamido-1-indanone (2.5 g, 13.2 mmol), which was stirred at 100°C for 3.5 hours, and cooled to room temperature. The precipitated crude products were collected, which was washed with ethyl acetate to give the titled compound (2.73 g).

¹H NMR (DMSO-d₆) δ : 2.08 (3H, s), 3.13 (6H, s), 3.87 (2H, s), 7.31 (1H, s), 7.52 (2H, m), 7.86 (1H, s), 10.16 (1H, s).

Reference Example 48

N-[2-[(Dimethylamino)methyl]-2,3-dihydro-1H-inden-5-yl]acetamide

15



15

N-[2-[(E)-(Dimethylamino)methylidene]-1-oxo-2,3-dihydro-1H-inden-5-yl]acetamide (2.70 g, 12.3 mmol) obtained in Reference Example 47 and 10% palladium-carbon (0.3 g) were added to a mixed solution of methanol (60 ml) and acetic acid (6 ml), which was stirred at 40°C under a hydrogen atmosphere for 1 day. After the catalyst was filtered, the filtrate was distilled out under reduced pressure. 1N hydrochloric acid (15 ml) was added to the reaction mixture, which was washed with ethyl acetate.

20

Then, potassium carbonate was added to the mixture, and extraction was conducted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, dried using anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was purified using aluminum column chromatography (development solvent: ethyl acetate) to give the titled

30

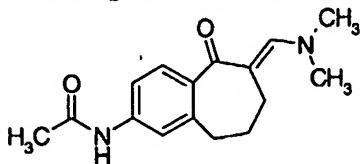
compound.

^1H NMR (CDCl_3) δ : 2.15 (3H, s), 2.25 (6H, s), 2.28 (2H, m), 2.61 (3H, m), 3.02 (2H, m), 7.11 (2H, m), 7.26 (1H, s), 7.39 (1H, s).

5

Reference Example 49

N-[6-[(E)-(Dimethylamino)methylidene]-5-oxo-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-2-yl]acetamide



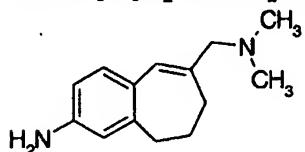
10 The titled compound was obtained by carrying out the same operation as in Reference Example 47, using N-(5-oxo-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-2-yl)acetamide.

15 $^1\text{H-NMR}$ (CDCl_3) δ : 1.78-1.90 (2H, m), 2.17 (3H, s), 2.34 (2H, t, $J = 6.6$ Hz), 2.74 (2H, t, $J = 6.8$ Hz), 3.11 (6H, s), 7.21 (1H, d, $J = 8.1$ Hz), 7.48-7.63 (3H, m), 7.73 (1H, s).

Melting point: 177 - 180°C (crystallization solvent: ethyl acetate-diethyl ether)

20 **Reference Example 50**

8-[(Dimethylamino)methyl]-6,7-dihydro-5H-benzo[a]cyclohepten-3-amine



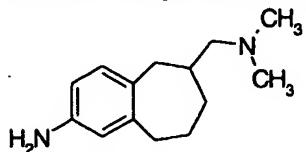
25 The titled compound was obtained as an oily substance by carrying out the same operation as in Example 41-2), using N-[6-[(E)-(dimethylamino)methylidene]-5-oxo-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-2-yl]acetamide obtained in Reference Example 49.

30 $^1\text{H-NMR}$ (CDCl_3) δ : 1.90-2.01 (2H, m), 2.22 (6H, s), 2.35 (2H, t, $J = 6.3$ Hz), 2.72 (2H, t, $J = 5.4$ Hz), 2.91 (2H, s), 3.7

(2H, br, NH₂), 6.28 (1H, s), 6.40-6.50 (2H, m), 6.94 (1H, d, J = 7.8 Hz).

Reference Example 51

5 6-[(Dimethylamino)methyl]-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-2-amine



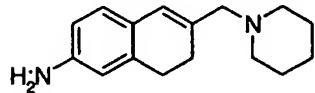
The titled compound was obtained as an oily substance, by carrying out the same operation as in Reference Example 10 48, using 8-[(dimethylamino)methyl]-6,7-dihydro-5H-benzo[a]cyclohepten-3-amine.

¹H-NMR (CDCl₃) δ : 1.30-1.63 (3H, m), 1.65-2.22 (10H, m), 2.44-2.80 (4H, m), 3.5 (2H, br, NH₂), 6.35-6.48 (2H, m), 6.92 (1H, d, J = 7.8 Hz).

15

Reference Example 52

6-(1-Piperidinylmethyl)-7,8-dihydro-2-naphthalenamine



1) A mixture of 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone (11 g) obtained in Example 41-1) and piperidine (100 ml) was refluxed with heating for 24 hours. After excess piperidine was distilled out under reduced pressure, the resulting residue was crystallized using tetrahydrofuran-isopropyl ether to give 6-acetamido-2-(1-piperidinylmethylidene)-1-tetralone (7 g) as a light yellow powder.

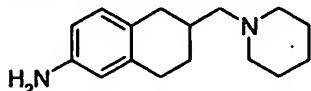
2) The titled compound was obtained as an amorphous powder by carrying out the same operations as in Example 41-2), using 6-acetamido-2-(1-piperidinylmethylidene)-1-tetralone obtained in above 1).

¹H NMR (CDCl₃) δ : 1.44-1.57 (6H, m), 2.25-2.34 (6H, m), 2.72 (2H, t, J=8.0 Hz), 2.98 (2H, s), 3.59 (2H, s), 6.23 (1H,

s), 6.45-6.47 (2H, m), 6.81 (1H, d, J=8.7 Hz).

Reference Example 53

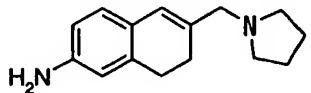
5 **6-(1-Piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine**



The titled compound was obtained as an amorphous powder by carrying out the same operations as in Reference Example 48, using 6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 52.
10 ¹H NMR (CDCl₃) δ: 1.25-2.82 (19H, m), 3.36 (2H, bs), 6.44-6.49 (2H, m), 6.88 (1H, d, J=8.1 Hz).

Reference Example 54

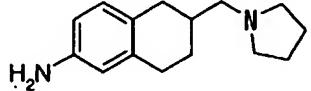
15 **6-(1-Pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine**



The titled compound was obtained as an amorphous powder by carrying out the same operations as in Reference Example 52, using 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone obtained in Example 41-1).
20 ¹H NMR (CDCl₃) δ: 1.76-1.80 (4H, m), 2.30 (2H, t, J = 7.8 Hz), 2.47-2.49 (4H, m), 2.74 (2H, t, J = 7.8 Hz), 3.13 (2H, s), 3.59 (2H, brs), 6.26 (1H, s), 6.45-6.47 (2H, m), 25 6.82 (1H, d, J = 8.6Hz).

Reference Example 55

6-(1-Pyrrolidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine



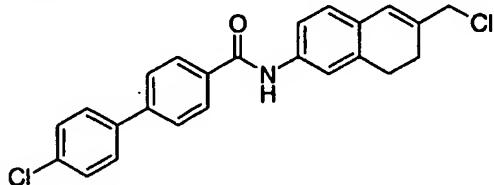
30 The titled compound was obtained as an amorphous

powder by carrying out the same operations as in Reference Example 48, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H NMR (CDCl₃) δ: 1.45-1.90 (1H, m), 1.55-2.80 (16H, m), 5 3.48 (2H, brs), 6.44 (1H, s), 6.47 (2H, d, J = 8.1 Hz), 6.88 (2H, d, J = 8.1 Hz).

Reference Example 56

10 **4'-Chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl] [1,1'-biphenyl]-4-carboxamide**

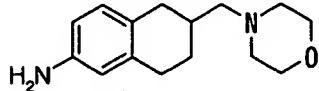


After 1-chloroethyl chloroformate (0.23 ml) was added to tetrahydrofuran solution (30 ml) of 4'-chloro-N-[6-(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide (750 mg) at 15 -78°C, the temperature of the solution was raised to room temperature over 30 minutes. The solvent was distilled out under reduced pressure. The resulting residue was crystallized using tetrahydrofuran-n-hexane to give the 20 titled compound (600 mg).

Melting point: 179 - 181°C (crystallization solvent:
tetrahydrofuran-n-hexane)

Reference Example 57

25 **6-(4-Morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine**



The titled compound was obtained as an amorphous powder by carrying out, in order, the same operations as 30 in Reference Example 52 and Reference Example 48, using 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-

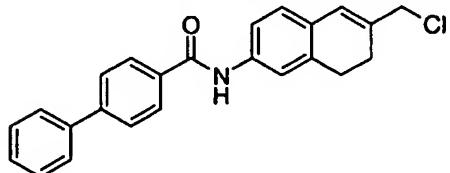
tetralone obtained in Example 41-1).

¹H NMR (CDCl₃) δ : 1.22-1.41 (1H, m), 1.80-1.82 (2H, m), 2.22-2.34 (10H, m), 3.50 (2H, s), 3.69-3.72 (1H, m), 6.40 (1H, s), 6.44 (1H, d, J = 8.1 Hz), 6.85 (1H, d, J = 8.1 Hz).

5

Reference Example 58

N-[6-(Chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

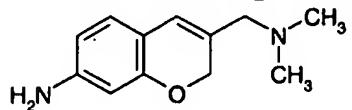


10 The titled compound was obtained by carrying out the same operations as in Reference Example 56, using N-[6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Example 47.

15 Melting point: 163 - 165°C (crystallization solvent: tetrahydofuran-n-hexane)

Reference Example 59

3-[(N,N-Dimethylamino)methyl]-2H-chromen-7-amine

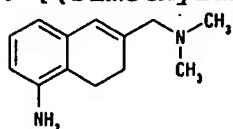


20 The titled compound was obtained by carrying out, in order, the same operations as in Examples 41-1) and 41-2), using 7-acetylamino-3,4-dihydrochromen-4-on.

25 ¹H-NMR (CDCl₃) δ : 2.20 (6H, s), 2.94 (2H, s), 3.66 (2H, brs), 4.71 (2H, s), 6.16-6.21 (2H, m), 6.76 (1H, d, J = 7.8 Hz).

Reference Example 60

6-[(Dimethylamino)methyl]-7,8-dihydro-1-naphthalenamine



1) Methyl 4-(2-aminophenyl)butanoate hydrochloride (7.20 g, 0.037 mol) synthesized by a known method by documents (*Synthetic communications*, 26(18), 3443 (1996)) and triethylamine (5.06 g, 0.05 mol) were dissolved in 5 tetrahydrofuran (60ml). Acetyl chloride (3.51 g, 0.045 mol) was added dropwise to the mixture, which was stirred at room temperature for 30 minutes. Ethyl acetate and 1N hydrochloric acid were added to the reaction mixture, and extraction was conducted. The organic layer was washed 10 with water, concentrated and dried. A mixed solution of ethyl acetate - n-hexane (1 : 1) was added to the residue.

The crystallized product was collected by filtration, to give methyl 4-(2-acetylaminophenyl)butanoate (6.40g) as a white powder.

15 $^1\text{H-NMR}$ (CDCl_3) δ : 1.77-1.86 (2H, m), 2.29 (3H, s), 2.41-2.45 (2H, m), 2.59-2.62 (2H, m), 3.74 (3H, s), 7.03 (1H, t, $J=7.3$ Hz), 7.11-7.12 (1H, m), 7.22 (1H, t, $J=7.3$ Hz), 8.08 (1H, d, $J=8.1$ Hz), 8.33 (1H, s).

20 2) Polyphosphoric acid (100g) was heated at 130°C, then methyl 4-(2-acetylaminophenyl)butanoate (6.40g, 0.027mol) obtained in 1) was added under stirring. After stirring for 1 hour, the reaction mixture was poured into ice water, and ethyl acetate and water were added, then extraction was conducted by adding water. The organic 25 layer was washed with saturated sodium hydrogen carbonate solution and aqueous sodium chloride solution, and concentrated. A mixed solution of ethyl acetate - n-hexane (1:1) was added to the residue, and the crystallized product was collected by filtration, to give 5-acetylamino-1-tetralone (2.80g) as a white powder.

30 $^1\text{H-NMR}$ (CDCl_3) δ : 2.10-2.19 (2H, m), 2.24 (3H, s), 2.66 (2H, t, $J=6.3$ Hz), 2.84 (2H, t, $J=5.7$ Hz), 7.06 (1H, brs), 7.34 (1H, t, $J=7.5$ Hz), 7.82(1H, d, $J=7.5$ Hz), 7.95 (1H, d, $J=7.5$ Hz).

35 3) 5-Acetylamino-1-tetralone (0.6g, 3.0 mmol) obtained was dissolved in dimethylformamide dimethylacetal

(20ml), which was refluxed with heating for 4 hours. The crystallized product was collected by filtration, which was washed with ethyl acetate, to give 5-acetylaminoo-2-(dimethylamino)methylidene-1-tetralone (0.58g) as a yellow powder.

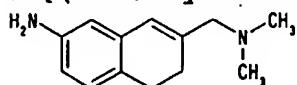
5 $^1\text{H-NMR}$ (CDCl_3) δ : 2.21 (3H, s), 2.68-2.72 (2H, m), 2.86-2.90 (2H, m), 3.11 (6H, s), 7.26-7.31 (2H, m), 7.62 (1H, m), 7.69 (1H, s), 7.92 (1H, m).

10 4) Sodium triacetoxyhydroborate (424 mg, 2.0 mmol) was dissolved in a mixed solution of ethyl acetate (5ml) and tetrahydrofuran (1ml) under ice-cooling. 5-Acetylaminoo-2-dimethylaminomethylidene-1-tetralone (129 mg, 0.5 mmol) obtained in 3) was added to the mixture, which was stirred for 15 minutes. The reaction mixture was 15 concentrated, and methanol (10ml) was added to the residue, and sodium borohydride (38 mg, 1 mmol) was added under ice-cooling. After stirring for 1 hour, the reaction mixture was concentrated. 5N Hydrochloric acid and ethyl acetate were added to the residue, and extraction was 20 conducted. The water layer was refluxed with heating for 2 hours. 4N sodium hydroxide solution and ethyl acetate were added to the reaction mixture, and extraction was conducted. The organic layer was washed with water, and concentrated. The residue was purified by alumina column 25 chromatography (development solvent: ethyl acetate : n-hexane=1:1), to give the titled compound (80 mg) as a colorless oily substance.

30 $^1\text{H-NMR}$ (CDCl_3) δ : 2.24(6H, s), 2.37(2H, t, $J=8.1$ Hz), 2.63(2H, t, $J=8.1$ Hz), 2.97(2H, s), 3.58(2H, brs), 6.29(1H, s,), 6.53(1H, d, $J=8.1$ Hz), 6.57 (1H, d, $J=8.1$ Hz), 6.97(1H, t, $J=8.1$ Hz).

Reference Example 61

7-[(Dimethylamino)methyl]-5,6-dihydro-2-naphthalenamine



1) 7-Nitro-1-tetralone (8.32 g, 0.044 mol) and concentrated hydrochloric acid (24 ml, 0.29 mol) were dissolved in methanol (100 ml), and an iron powder (7.30 g, 0.13 mol) was gradually added over 1 hour. After 5 stirring for 1 hour, the reaction mixture was concentrated.

4N Sodium hydroxide solution and ethyl acetate were added to the residue, and extraction was conducted. The organic layer was dried, and concentrated. Tetrahydrofuran (100 ml) and triethylamine (5.05 g, 0.05 mol) was added to the 10 residue. Further, acetyl chloride (3.92 g, 0.05 mol) was added under ice-cooling. After stirring for 30 minutes, ethyl acetate and 1N hydrochloric acid were added, and extraction was conducted. The organic layer was concentrated, and the residue was purified with silica gel 15 column chromatography (development solvent: ethyl acetate), to give 7-acetylamino-1-tetralone (7.52 g) as a white powder.

¹H-NMR (CDCl₃) δ : 2.09-2.18 (2H, m), 2.21(3H, s), 2.65 (2H, t, J=6.3 Hz), 2.94 (2H, t, J=6.3 Hz), 7.24 (1H, d, J=8.4 Hz), 7.82 (1H, s), 7.98 (1H, brs), 8.15 (1H, d, J=7.5 Hz).

2) 7-Acetylamino-2-[(dimethylamino)methylidene]-1-tetralone (2.95 g) was obtained as a white powder by the same method as in Reference Example 60-3), using 7-acetylamino-1-tetralone (3.00 g, 0.0148 mol) obtained in 25 1).

¹H-NMR (CDCl₃) δ : 2.17 (3H, s), 2.78-2.82 (2H, m), 2.88-2.93 (2H, m), 3.14 (6H, s), 7.14 (1H, d, J=8.1 Hz), 7.74 (1H, s), 7.76 (1H, s), 8.09-8.12 (1H, m), 8.24 (1H, s).

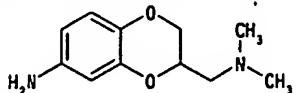
3) The titled compound (300 mg) was obtained as a 30 colorless oily substance by the same method as in Reference Example 60-4), using 7-acetylamino-2-[(dimethylamino)methylidene]-1-tetralone (628 mg, 2.43 mmol) obtained in 2).

¹H-NMR (CDCl₃) δ : 2.23 (6H, s); 2.29 (2H, t, J=8.4 Hz), 2.71 (2H, t, J=8.4 Hz), 2.97 (2H, s), 3.52 (2H, brs), 6.24 (1H, s,), 6.41 (1H, s,), 6.46 (1H, d, J=7.8 Hz), 6.90 (1H, d,

J=7.8 Hz).

Reference Example 62

N,N-Dimethyl-N-[(7-amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine



1) 1,2-Dihydroxy-4-nitrobenzene (5.00 g, 0.032 mol), potassium carbonate (9.67 g, 0.07 mol) and epibromohydrin (5.30 g, 0.039 mol) were dissolved in dimethylformamide (100ml), which was stirred at 100°C for 1 hour. Water was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water, and concentrated. The residue was purified by alumina column chromatography (development solvent : ethyl acetate). The eluent was washed with a mixed solution of ethyl acetate - n-hexane (1:1), to give (7-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methanol (3.31 g) as a white powder.

¹H-NMR (CDCl₃) δ : 1.95-1.99 (1H, m), 3.89-3.97 (2H, m), 4.19-4.29 (2H, m), 4.41-4.45 (1H, m), 6.96 (1H, d, J=8.6 Hz), 7.78-7.81 (2H, m).

2) (7-Nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methanol (1.00 g, 4.74 mmol) obtained in 1) and triethylamine (719 mg, 7.10 mmol) were dissolved in dimethylformamide (30 ml), and methanesulfonyl chloride (651 mg, 5.68 mmol) was added, which was stirred at room temperature for 30 minutes. Then, an aqueous dimethylamine solution was added and stirred at 60°C for 5 hours. Water was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water, and concentrated. The residue was purified by alumina column chromatography (development solvent : ethyl acetate : n-hexane = 3:7), to give N,N-dimethyl-N-[(7-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine (802 mg) as a colorless oily substance.

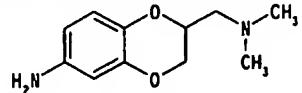
¹H-NMR (CDCl₃) δ : 2.34 (6H, s), 2.50-2.68 (2H, m), 4.02-4.09 (2H, m), 4.30-4.36 (1H, m), 4.39-4.44 (2H, m), 6.94 (1H, d, J=8.9Hz), 7.76-7.84 (2H, m).

3) N,N-Dimethyl-N-[(7-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine (802 mg, 3.37 mmol) obtained in 2) and concentrated hydrochloric acid (3 ml) was dissolved in methanol (10 ml), and an iron powder (0.80 g, 14 mmol) was quietly added over 1 hour. After stirring for 1 hour, the reaction mixture was concentrated. 4N Sodium hydroxide solution and ethyl acetate were added to the residue, and extraction was conducted. The organic layer was dried, and concentrated. The residue was purified by silica gel column chromatography (development solvent: ethyl acetate - n-hexane = 3:7), to give the titled compound (514 mg) as a colorless oily substance.

¹H-NMR (CDCl₃) δ : 2.32 (6H, s), 2.43-2.64 (2H, m), 3.40 (2H, s), 3.86-3.93 (1H, m), 4.19-4.27 (2H, m), 6.18-6.22 (1H, m), 6.29 (1H, s), 6.67 (1H, d, J=8.7 Hz).

20 Reference Example 63

N,N-Dimethyl-N-[(6-amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine



1) 1,2-Dihydroxy-4-nitrobenzene (4.65 g, 0.030 mol), 25 potassium carbonate (8.71 g, 0.063 mol) and methoxymethyl chloride (2.42 g, 0.030 mol) were dissolved in dimethylformamide (50 ml), which was stirred at 40°C for 30 minutes. Epibromohydrin (7.20 g, 0.045 mol) was added to the mixture, which was stirred at 60°C for 80 minutes. 30 Then water was added, and extraction was conducted using ethyl acetate. The organic layer was washed with water, and concentrated. The residue was purified by alumina column chromatography (development solvent: ethyl acetate - n-hexane = 1:4), to give 2-[(2-(methoxymethoxy)-5-nitrophenoxy)methyl]oxirane (2.61 g) as a white powder.

¹H-NMR (CDCl₃) δ : 2.79-2.81 (1H, m), 2.93-2.96 (1H, m), 3.41 (1H, m), 3.53 (3H, s), 4.01-4.07 (1H, m), 4.40-4.45 (1H, m), 5.32 (2H, s), 7.22 (1H, d, J=9.0 Hz), 7.82-7.91 (2H, m).

5 2) 2-[2-(Methoxymethoxy)-5-nitrophenoxy]methyl]oxirane (4.00 g, 0.016 mol) obtained in 1) was dissolved in methanol (50 ml), and 10% hydrochloric acid-methanol solution (10 ml) was added, which was stirred at room temperature for 30 minutes. The 10 solvent was concentrated, and methanol (30 ml) and potassium carbonate (6.50 g, 0.047 mol) were added to the residue, which was stirred at 60°C for 1 hour. The solvent was concentrated, water was added, and extraction was conducted using ethyl acetate. The organic layer was 15 washed with water, and concentrated. The residue was purified by alumina column chromatography (development solvent; ethyl acetate), to give (6-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methanol (2.12 g) as a white powder.
¹H-NMR (CDCl₃) δ : 1.90-1.94 (1H, m), 3.89-3.97 (2H, m), 4.19-4.28 (2H, m), 4.41-4.45 (1H, m), 6.97 (1H, d, J=8.6 Hz), 7.78-7.82 (2H, m).

20 3) N,N-Dimethyl-N-[(6-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine (910 mg) was obtained as a colorless oily substance, by the same method as in Reference 25 Example 62-2), using (6-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methanol (1.00 g, 4.74 mmol) obtained in 2).

25 ¹H-NMR (CDCl₃) δ : 2.35 (6H, s), 2.52-2.70 (2H, m), 3.98-4.05 (2H, m), 4.35-4.39 (3H, m), 6.95-6.98 (1H, m), 7.77-7.80 (2H, m).

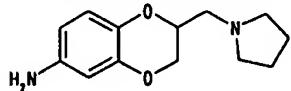
30 4) The titled compound (750 mg) was obtained as a colorless oily substance, by the same method as in Reference Example 62-3), using N,N-dimethyl-N-[(6-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine (910 mg, 3.82 mmol) obtained in 3).

35 ¹H-NMR (CDCl₃) δ : 2.32 (6H, s), 2.43-2.64 (2H, m), 3.40 (2H,

s), 3.86-3.92 (1H, m), 4.13-4.27 (2H, m), 6.19-6.28 (2H, m), 6.67-6.70 (1H, m).

Reference Example 64

5 1-[(6-Amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]pyrrolidine

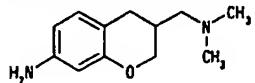


10 1) 1-[(6-Nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]pyrrolidine (1.30 g) was obtained as a colorless oily substance, by the same method as in Reference Example 62-2), using (6-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methanol (1.12 g, 5.30 mmol) and pyrrolidine (10 ml).
¹H-NMR (CDCl₃) δ : 1.79-1.83 (4H, m), 2.60-2.62 (4H, m), 2.78 (2H, d, J=5.9 Hz), 4.00-4.07 (1H, m), 4.38-4.42 (2H, m), 15 6.95-6.98 (1H, m), 7.76-7.80 (2H, m).

15 2) The titled compound (1.03 g) was obtained as a colorless oily substance, by the same method as in Reference Example 62-3), using 1-[(6-nitro-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]pyrrolidine (1.30 g, 4.92 mmol).
¹H-NMR (CDCl₃) δ : 1.74-1.83 (4H, m), 2.54-2.63 (4H, m), 2.69-2.72 (2H, m), 3.40 (2H, s), 3.91-3.97 (1H, m), 4.18-4.30 (2H, m), 6.18-6.25 (2H, m), 6.70 (1H, d, J=8.4 Hz).

25 **Reference Example 65**

N-[(7-Amino-3,4-dihydro-2H-chromen-3-yl)methyl]-N,N-dimethylamine



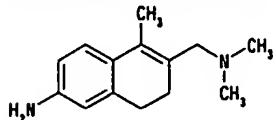
30 3-[(N,N-Dimethylamino)methyl]-2H-chromen-7-amine (150 mg, 0.73 mmol) obtained in Reference Example 59, 1N hydrochloric acid (0.5 ml) and 10% palladium carbon (40 mg) was dissolved in methanol (5 ml), and catalytic hydrogenation was conducted under normal temperature and normal pressure. After a catalyst was filtered out, the

filtrate was concentrated, and the residue was purified by alumina column chromatography (development solvent; ethyl acetate : n-hexane = 3:7), to give the titled compound (15 mg) as a colorless oily substance.

5 ¹H-NMR (CDCl₃) δ: 2.20-2.24 (3H, m), 2.24(6H, m), 2.30-2.40 (1H, m), 2.75-2.80 (1H, m), 3.60 (1H, m), 3.75-3.80 (2H, m), 4.20-4.25 (1H, m), 6.20 (1H, m), 6.21-6.25 (1H, m), 6.82 (1H, d, J=7.8 Hz).

10 Reference Example 66

6-[(Dimethylamino)methyl]-5-methyl-7,8-dihydro-2-naphthalenamine



15 1) 6-Acetylaminio-1-tetralone (5.5 g, 0.027 mol) and dimethylmethylenammonium chloride (6.3 g, 0.068 mol) were dissolved in a mixed solution of acetonitrile (100 ml) and tetrahydrofuran (100 ml), which was stirred for 48 hours.

20 The crystallized product was collected by filtration, washed with tetrahydrofuran, and dissolved in ethyl acetate. 0.5N Sodium hydroxide solution was added to the solution for liquid separation. The organic layer was concentrated, to give 6-acetylaminio-2-[(dimethylamino)methyl]-1-tetralone (4.48 g) as a colorless oily substance.

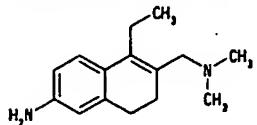
25 2) 6-Acetylaminio-2-[(dimethylamino)methyl]-1-tetralone (260 mg, 1.00 mmol) obtained was dissolved in tetrahydrofuran (10 ml). 1M Methyl magnesium bromide - tetrahydrofuran solution (3 ml)(3.00 mmol) was added to the solution under ice-cooling, which was stirred at room 30 temperature for 16 hours. Aqueous ammonium chloride solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was concentrated, and 5N hydrochloric acid and ethyl acetate were added to the residue for liquid separation.

Concentrated hydrochloric acid was added to the water layer, which was refluxed for 4 hours. The reaction mixture was concentrated, and 1N sodium hydroxide solution and ethyl acetate were added to the residue and extraction was conducted. The organic layer was concentrated, and the residue was purified by alumina column chromatography (development solvent; ethyl acetate : n-hexane = 3:7), to give the titled compound (83 mg) as a colorless oily substance.

¹H-NMR (CDCl₃) δ : 2.04 (3H, s), 2.24 (6H, s), 2.28 (2H, t, J=7.4 Hz), 2.66 (2H, t, J=7.4 Hz), 3.04 (2H, s), 3.62 (2H, s), 6.49 (1H, s), 6.51-6.55 (1H, m), 7.10 (1H, d, J=8.1 Hz).

15 Reference Example 67

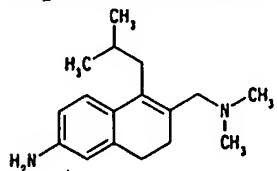
6-[(Dimethylamino)methyl]-5-ethyl-7,8-dihydro-2-naphthalenamine



The titled compound was obtained as a colorless oily substance by the same manner as in Reference Example 66-2), using 6-acetylmino-2-(dimethylamino)methyl-1-tetralone obtained in Reference Example 66-1) and ethyl magnesium bromide.

¹H-NMR (CDCl₃) δ : 1.06 (3H, t, J=7.5 Hz), 2.24 (6H, s), 2.27 (2H, m), 2.52-2.66 (4H, m), 3.04 (2H, s), 3.61 (2H, s), 6.51 (1H, s), 6.51-6.55 (1H, m), 7.11 (1H, d, J=8.1 Hz).

Reference Example 68
6-[(Dimethylamino)methyl]-5-isobutyl-7,8-dihydro-2-naphthalenamine



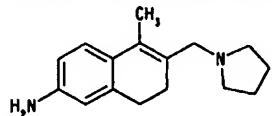
The titled compound was obtained as a colorless oily substance by the same manner as in Reference Example 66-2), using 6-acetylamino-2-[(dimethylamino)methyl]-1-tetralone obtained in Reference Example 66-1) and isobutyl 5 magnesium bromide.

¹H-NMR (CDCl₃) δ : 0.88 (6H, d, J=6.7 Hz), 1.73-1.79 (1H, m), 2.21 (6H, s), 2.28 (2H, t, J=7.0 Hz), 2.44 (2H, d, J=7.3 Hz), 2.63 (2H, t, J=7.0 Hz), 3.09 (2H, s), 3.60 (2H, s), 6.49 (1H, s), 6.51-6.53 (1H, m), 7.08 (1H, d, J=7.8 Hz).

10

Reference Example 69

5-Methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine



15 1) 6-Acetylamino-2-[(dimethylamino)methylidene]-1-tetralone (4.90 g, 0.017 mol) obtained in Example 41-1) was suspended in pyrrolidine (25 ml), which was refluxed with heating for 2 hours. The crystallized product was collected by filtration, washed with a mixed solution of 20 ethyl acetate and n-hexane (1:1), to give 6-acetylamino-2-(1-pyrrolidinylmethylidene)-1-tetralone (5.03 g) as yellow crystals.

17 ¹H-NMR (CDCl₃) δ : 1.75-2.00 (4H, m), 2.19 (3H, s), 2.70-3.00 (4H, m), 3.50-3.70 (4H, m), 7.20-7.25 (1H, m), 7.67 (1H, s), 7.70-7.90 (2H, m), 7.97 (1H, d, J=8.4 Hz).

22 2) Sodium triacetoxyhydroborate (3.18 g, 0.015 mol) was dissolved in a mixed solution of ethyl acetate (50 ml) and tetrahydrofuran (12.5 ml) under ice-cooling, and 6-acetylamino-2-(1-pyrrolidinylmethylidene)-1-tetralone (2.84 g, 0.01mol) obtained in 1) was added. After stirring for 1 hour, the reaction mixture was concentrated. 1N Sodium hydroxide solution and ethyl acetate were added to the residue, which was stirred. The crystallized product was collected by filtration, washed with a mixed solution

of ethyl acetate and n-hexane (1:1), to give 6-acetylamino-2-(1-pyrrolidinylmethyl)-1-tetralone (2.65 g) as a white powder.

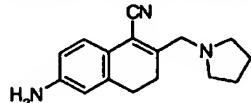
¹H-NMR (CDCl₃) δ : 1.78 (4H, m), 1.90-2.02 (1H, m), 2.20 (3H, s), 2.35-2.98 (10H, m), 7.20-7.23 (1H, m), 7.57 (1H, s), 7.66 (1H, m), 7.97 (1H, d, J=8.4 Hz).

3) The titled compound was obtained by the same manner as in Reference Example 66-2), using 6-acetylamino-2-(1-pyrrolidinylmethyl)-1-tetralone obtained in 2).

10 ¹H-NMR (CDCl₃) δ : 1.73-1.79 (4H, m), 2.04 (3H, s), 2.31 (2H, t, J=7.4 Hz), 2.49-2.54 (4H, m), 2.65 (2H, t, J=7.8 Hz), 3.24 (2H, s), 3.60 (2H, brs), 6.48-6.54 (2H, m), 7.09 (1H, d, J=8.1 Hz).

15 Reference Example 70

6-Amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-1-naphthalenecarbonitrile



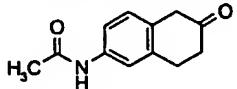
Trimethylsilylnitrile (1.02 ml, 7.68 mmol) and zinc iodide (22 mg, 0.0698 mmol) were added to dichloroethane solution (9 ml) of 6-acetylamino-2-(1-pyrrolidinylmethyl)-1-tetralone (1.00 g, 3.49 mmol) obtained in Reference Example 69-2), which was stirred at room temperature for 2 days. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the obtained oily substance, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was purified by alumina column chromatography (development solvent; ethyl acetate), to give trimethylsilylcyanohydrin form (1.21 g) as an oily substance. 2.5N Hydrochloric acid was added to the oily substance (978 mg, 2.73 mmol), which was stirred at 100°C for 1.5 hours. The aqueous solution obtained was

washed with ethyl acetate. Potassium carbonate was added to the water layer to make it alkaline, and extraction was conducted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, dried over
5 anhydrous sodium sulfate, and the solvent was distilled out under reduced pressure. The resulting oily substance was purified by alumina column chromatography (development solvent : hexane : ethyl acetate = 5:1), to give the titled compound (358 mg).

10 ^1H NMR (CDCl_3) δ : 1.80 (4H, m), 2.56 (6H, m), 3.73 (2H, m),
3.50 (2H, s), 3.77 (2H, br), 6.46 (1H, s), 6.55 (1H, d, J = 8.1 Hz), 7.26 (1H, d, J = 8.1 Hz).

Reference Example 71

15 6-Acetamido-2-tetralone



1) Sodium borohydride (931 mg, 24.6 mmol) was added to a methanol solution (60 ml) of 6-acetamido-1-tetralone (5.00 g, 24.6 mmol) under ice-cooling, which was stirred
20 at room temperature for 1 hour. Ethyl acetate was added to the reaction mixture, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then, the solvent was distilled out under reduced pressure. p-Toluenesulfonic acid (468 mg,
25 2.46 mmol) and toluene (120 ml) were added to the obtained alcohol form (5.05 g, 24.6 mmol), which was stirred at 100 °C for 1 hour. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with saturated aqueous sodium chloride solution,
30 dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by silica gel column chromatography (development solvent; hexane:ethyl acetate = 1:1), and powdered with hexane to give N-(7,8-dihydro-2-naphthalenyl)acetamide (3.17 g).

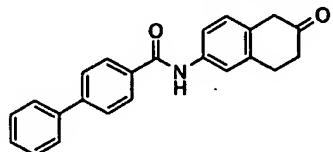
¹H NMR (CDCl₃) δ : 2.16 (3H, s), 2.29 (2H, m), 2.28 (2H, m), 5.97 (1H, m), 6.42 (2H, d, J=9.6 Hz), 6.97 (1H, d, J=8.1 Hz), 7.14 (1H, br), 7.20 (1H, .m), 7.32 (1H, s).

2) m-Chloroperbenzoic acid (5.13 g, 20.8 mmol) was
5 added to a chloroform solution (80 ml) of N-(7,8-dihydro-2-naphthalenyl)acetamide (3.00 g, 16.0 mmol) obtained in 1) under ice-cooling, which was stirred at room temperature for 2 hours. Ethyl acetate was added to the reaction mixture, which was washed with saturated sodium
10 hydrogencarbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by alumina B column chromatography (development solvent; hexane : ethyl acetate = 1:1). 1N Sodium hydroxide solution (10.7 ml) was added to a methanol solution (100 ml) of the obtained
15 oily substance (3.20 g, 8.89 mmol) under ice-cooling, which was stirred at room temperature for 30 minutes. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by alumina B column chromatography (development
20 solvent; ethyl acetate : methanol = 10:1). p-Toluenesulfonic acid (50mg, 0.262 mmol) and toluene (26 ml) were added to the obtained diol (596 mg, 2.62 mmol), which was stirred at 120°C for 3 hours. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the
25 residue, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by silica gel column chromatography (development solvent; hexane : ethyl acetate = 1:3), and powdered with diisopropyl ether, to give the titled compound (231 mg).
30
35

¹H NMR (CDCl₃) δ : 2.18 (3H, s), 2.54 (2H, m), 3.04 (2H, m), 3.76 (2H, s), 7.06 (1H, d, J=8.1 Hz), 7.21 (1H, dd, J=8.1, 2.0 Hz), 7.31 (1H, br), 7.61 (1H, d, J=2.0 Hz).

5 Reference Example 72

N-(6-Oxo-5,6,7,8-tetrahydro-2-naphthalenyl)[1,1'-biphenyl]-4-carboxamide



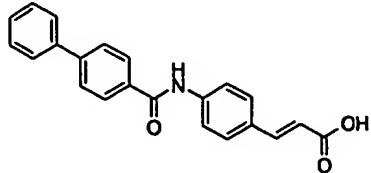
Concentrated hydrochloric acid (1.5 ml) was added to
10 6-acetamido-2-tetralone (20 mg, 0.098 mmol) obtained in
Reference Example 71, which was stirred at 100°C for 1 hour,
and the solvent was distilled out under reduced pressure.

Ethyl acetate was added to the residue, which was washed
with aqueous potassium carbonate solution and saturated
15 aqueous sodium chloride solution, dried over anhydrous
sodium sulfate, and then the solvent was distilled out under
reduced pressure. [1,1'-Biphenyl]-4-carbonyl chloride
(21.3 mg, 0.098 mmol) was added to a dimethylformamide
solution (0.5 ml) of the obtained oily substance and
20 triethylamine (0.014 ml, 0.098 mmol) under ice-cooling,
which was stirred at room temperature for 1 hour. Ethyl
acetate was added to the reaction mixture, which was washed
with 1N hydrochloric acid, aqueous potassium carbonate
solution and saturated aqueous sodium chloride solution,
25 dried over anhydrous sodium sulfate, and then the solvent
was distilled out under reduced pressure. The resulting
residue was purified by silica gel column chromatography
(development solvent;hexane:ethyl acetate = 1:1), to give
the titled compound (10 mg).

30 ¹H NMR (CDCl₃) δ : 2.56 (2H, t, J=6.6 Hz), 3.08 (2H, t, J=6.6
Hz), 3.57 (2H, s), 7.11 (1H, d, J=8.1 Hz), 7.43 (4H, m),
7.64 (2H, m), 7.72 (3H, m), 7.96 (3H, m).

Reference Example 73

(E)-3-[4-[(1,1'-biphenyl)-4-ylcarbonyl]amino]phenyl]-2-propenic acid

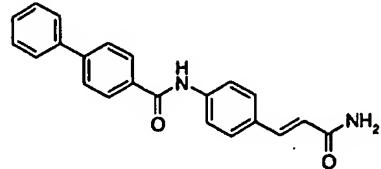


5 4-Phenylbenzoyl chloride (2.00 g, 9.23 mmol) was added to a mixed solution of 4-aminocinnamic acid (1.51 g, 9.23 mmol) and sodium hydrogen carbonate (2.33 g, 27.7 mmol) in water and diethyl ether under ice-cooling, which was stirred for 5 hours. After the reaction mixture was
10 separated, 5N hydrochloric acid was added to water layer, and the precipitated crude product was washed with water and ethyl acetate, to give the titled compound (1.34 g).
¹H NMR (DMSO-d₆) δ : 6.84 (1H, d, J = 16.0 Hz), 7.43-7.93 (12H, m), 8.09 (2H, d, J = 8.4 Hz), 10.51 (1H, s).

15

Reference Example 74

N-[4-[(E)-3-Amino-3-oxo-1-propenyl]phenyl][1,1'-biphenyl]-4-carboxyamide



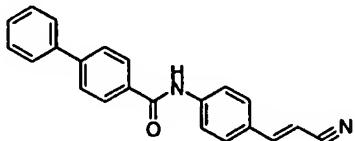
20 Chloro isobutylcarbonate (0.453 ml, 3.49 mmol) was added to a dimethylformamide suspension of (E)-3-[4-[(1,1'-biphenyl)-4-ylcarbonyl]amino]phenyl]-2-propionic acid (1.00 g, 2.91 mmol) obtained in Reference Example 73 and triethylamine (0.527 ml, 3.79 mmol) under ice-cooling, which was stirred for 30 minute. The solvent was distilled out under reduced pressure. Sodium hydrogencarbonate solution was added to the residue, and the precipitated crude product was washed with water and acetonitrile, to give the titled compound (936 mg).

25

¹H NMR (DMSO-d₆) δ : 6.56 (1H, d, J = 15.6 Hz), 7.05 (1H, br), 7.52 (7H, m), 7.86 (6H, m), 8.08 (2H, d, J = 7.6 Hz).

Reference Example 75

5 N-[4-[(E)-2-Cyanoethenyl]phenyl][1,1'-biphenyl]-4-carboxamide

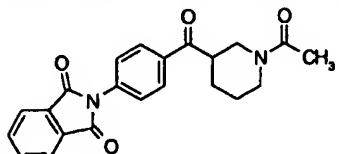


Cyanuric chloride (727 mg, 3.94 mmol) was added to a dimethylformamide suspension of (E)-3-[4-[(1,1'-biphenyl)-4-ylcarbonyl]amino]phenyl]-2-propenoic acid (900 mg, 2.63 mmol) obtained in Reference Example 74 at room temperature, which was stirred for 1 hour. After the solvent was distilled out under reduced pressure, the residue was dissolved in chloroform, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was stillled out under reduced pressure. The resulting residue was purified by silica gel column chromatography (development solvent; chloroform: ethyl acetate = 20:1), to give the titled compound (561 mg) as a colorless powder from diethyl ether.

¹H NMR (DMSO-d₆) δ : 6.37 (1H, d, J = 16.4 Hz), 7.43-7.51 (4H, m), 7.65-7.93 (8H, m), 8.08 (2H, d, J = 8.6 Hz).

25 Reference Example 76

2-[4-[(1-Acetyl-3-piperidinyl)carbonyl]phenyl]-1H-isoindol-1,3(2H)-dione



1) Thionyl chloride (2.12 ml, 32.1 mmol) was added to 30 fluorobenzene solution (20 ml) of 1-acetyl-3-

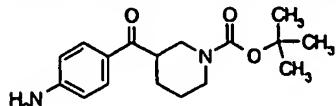
piperidinecarboxylic acid (5.00 g, 29.2 mmol) under ice-cooling, which was stirred at room temperature for 30 minutes. Aluminum chloride (9.74 g, 73.0 mmol) was added to the solution, which was stirred at 90°C for 1 hour. The 5 reaction mixture was poured in ice, and extraction was conducted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, saturated sodium hydrogencarbonate solution, and again saturated aqueous sodium chloride solution, dried over anhydrous 10 sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by silica gel column chromatography (development solvent; hexane:ethyl acetate = 1:1), to give (1-acetyl-3-piperidinyl)(4-fluorophenyl)methanone (4.93 g). 15 ^1H NMR (CDCl_3) δ : 1.61 (2H, m), 1.80 (2H, m), 2.11 and 2.15 (3H, s and s), 2.71 (1H, m), 3.11 and 3.42 (2H, m), 3.87 (1H, m), 4.53 and 4.83 (1H, m), 7.18 (2H, m), 8.02 (2H, m).

2) A dimethylformamide solution (50 ml) of (1-acetyl-3-piperidinyl)(4-fluorophenyl)methanone (4.92 g, 19.7 mmol) obtained in 1) and potassium phthalimide (3.66g, 19.7mmol) was stirred at 100°C for 12 hours under nitrogen atmosphere. The insoluble matters were filtered off, and the solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed 20 with 1N hydrochloric acid and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by silica gel column chromatography (development solvent; ethyl acetate), to give the titled compound (4.18 g) as a colorless powder from ethyl acetate - diisopropyl ether 25 (1:5).

^1H NMR (CDCl_3) δ : 1.66 (2H, m), 1.86 (2H, m), 2.13 and 2.15 (3H, s and s), 2.74 (1H, m), 3.11 and 3.43 (2H, m), 3.88 (1H, m), 4.54 and 4.85 (1H, m), 7.66 (2H, m), 7.82 (2H, m), 35 7.99 (2H, m), 8.10 (2H, m).

Reference Example 77

tert-Butyl 3-(4-aminobenzoyl)-1-piperidinecarboxylate



5 1) Concentrated hydrochloric acid (53 ml) was added to 2-[4-[(1-acetyl-3-piperidinyl)carbonyl]phenyl]-1H-isoindol-1,3(2H)-dione (4.00 g, 10.6 mmol) obtained in Reference Example 76, which was stirred at 100°C for 16 hours, and then insoluble matters were filtered off.

10 Potassium carbonate was added to the filtrate to make it alkaline, and extraction was conducted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure.

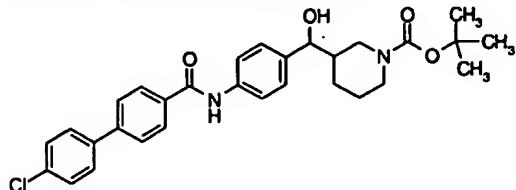
15 The resulting residue was powdered with diisopropyl ether, to give (4-aminophenyl)(3-piperidinyl)methanone (1.69 g). ¹H NMR (CD₃OD) δ : 1.59-1.85 (4H, m), 2.68-2.72 (2H, m), 3.30 (2H, m), 3.45 (1H, m), 6.62 (2H, m), 7.74 (2H, m).

20 2) t-Butyl dicarbonate (0.562 ml, 2.45 mmol) was added to a tetrahydrofuran solution (12 ml) of (4-aminophenyl)(3-piperidinyl)methanone (500 mg, 2.45 mmol) obtained in 1) under ice-cooling, which was stirred for 1.5 hours. Ethyl acetate was added to the reaction mixture, which was washed with saturated sodium hydrogen carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by silica gel column chromatography (development solvent; hexane:ethyl acetate = 1:1), to give the titled compound (831 mg).

25 ¹H NMR (CDCl₃) δ 1.47 (9H, s), 1.47-1.52 (2H, m), 1.67-1.74 (2H, m), 2.00 (1H, m), 2.72 (1H, m), 2.90 (1H, m), 3.32 (1H, m), 4.13 (3H, m), 6.66 (2H, d, J=8.4Hz), 7.84 (2H, d, J=8.4Hz).

Reference Example 78

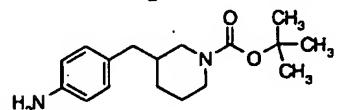
tert-Butyl 3-[[4-[(4'-chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]phenyl](hydroxy)methyl]-1-piperidinecarboxylate



tert-Butyl 3-[[4-[(4'-chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]benzoyl]-1-piperidinecarboxylate (506 mg, 0.975 mmol) obtained in Example 127-1) was dissolved
10 in a mixed solution of methanol and tetrahydrofuran (1:1) (10 ml). Sodium borohydride (73.8 mg, 1.95 mmol) was added to the solution under ice-cooling, which was stirred at room temperature for 1 hour. Ethyl acetate was added to the reaction mixture, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. Diisopropyl ether was added to the residue, to give the titled compound (488mg) as a colorless powder.
15
20 FABMS(pos) 521.2 [M+H]+

Reference Example 79

tert-Butyl 3-(4-aminobenzyl)-1-piperidinecarboxylate



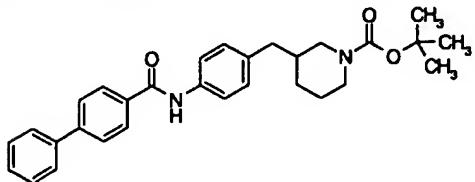
Sodium borohydride (433 mg; 11.5 mmol) was added to a methanol solution (25 ml) of tert-butyl 3-(4-aminobenzyl)-1-piperidinecarboxylate (1.74g, 5.73mmol) obtained in Reference Example 77 under ice-cooling, which was stirred at room temperature for 1 hour. Ethyl acetate was added to the reaction mixture, which was washed with saturated aqueous sodium chloride solution, dried over
25
30

anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by alumina B column chromatography (development solvent; ethyl acetate), to give an alcohol form. 1N hydrochloric acid (9.79 ml) and 10% palladium carbon (200 mg) were added to a methanol solution (300 ml) of the obtained alcohol form (1.00 g, 3.26 mmol), which was stirred for 16 hours under hydrogen atmosphere. The catalyst was filtered off, potassium carbonate was added to the filtrate to make it alkaline, and then the solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by silica gel column chromatography (development solvent; hexane - ethyl acetate = 1:1), to give the titled compound (813 mg).

¹H NMR (CDCl₃) δ: 1.46-1.76 (14H, m), 2.25-2.80 (2H, m), 3.14 (2H, m), 3.76 (4H, m), 6.64 (2H, m), 7.01 (2H, m).

Reference Example 80

tert-Butyl 3-[4-[(1,1'-biphenyl)-4-ylcarbonyl]amino]benzyl]-1-piperidinecarboxylate



The titled compound was obtained by carrying out the same operation as in Example 1, using tert-butyl 3-(4-aminobenzyl)-1-piperidinecarboxylate obtained in Reference Example 79 and [1,1'-biphenyl]-4-carboxylic acid.

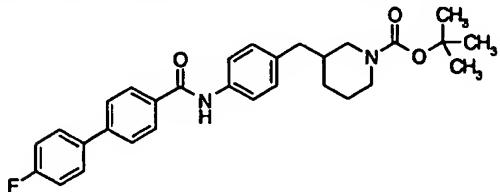
Elemental analysis for C₃₀H₃₄N₂O₃ · 0.5H₂O

Calcd.: C, 75.13; H, 7.36; N, 5.84.

Found : C, 74.83; H, 7.25; N, 5.65.
 Melting point: 135 - 137°C

Reference Example 81

5 tert-Butyl 3-[4-[(4'-fluoro[1,1'-biphenyl]-4-yl)carbonyl]amino]benzyl]-1-piperidinecarboxylate



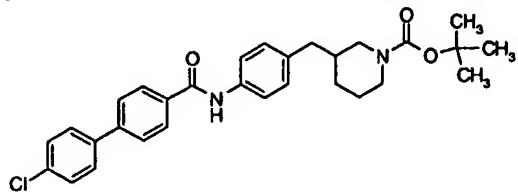
The titled compound was obtained by carrying out the same operation as in Example 1, using tert-butyl 3-(4-aminobenzyl)-1-piperidinecarboxylate obtained in Reference Example 80 and 4'-fluoro[1,1'-biphenyl]-4-carboxylic acid.

Elemental analysis for $C_{30}H_{33}FN_2O_3 \cdot 0.5H_2O$
 Calcd.: C, 72.41; H, 6.89; N, 5.63.
 15 Found: C, 72.30; H, 7.07; N, 5.60.

Melting point: 138 - 141°C

Reference Example 82

20 tert-Butyl 3-[4-[(4'-chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]benzyl]-1-piperidinecarboxylate



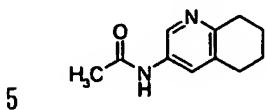
The titled compound was obtained by carrying out the same operation as in Example 1, using tert-butyl 3-(4-aminobenzyl)-1-piperidinecarboxylate obtained in Reference Example 80 and 4'-chloro[1,1'-biphenyl]-4-carboxylic acid.

Elemental analysis for $C_{30}H_{33}ClN_2O_3 \cdot 0.5H_2O$
 Calcd.: C, 70.09; H, 6.67; N, 5.45.
 Found: C, 70.29; H, 6.50; N, 5.38.

Melting point: 173 - 176°C

Reference Example 83

N-(5,6,7,8-Tetrahydro-3-quinolinyl)acetamide



1) Fuming nitric acid (100 ml) was added dropwise to concentrated sulfuric acid solution (200 ml) of 1-methyl-2-pyridone (20.7 g, 190 mmol) at 100°C, which was stirred for 16 hours. The reaction mixture was poured in ice. The 10 resulting precipitate was collected, which was washed with water, to give 1-methyl-3,5-dinitro-2(1H)-pyridinone (3.0 g).

¹H NMR (DMSO-d₆) δ: 3.68 (3H, s), 9.01 (1H, d, J=3.0 Hz), 9.61 (1H, d, J=3.0 Hz).

15 2) 1N Methanolic ammonia solution (300 ml) of 1-methyl-3,5-dinitro-2(1H)-pyridinone (3.00g, 15.1mmol) obtained in 1) and 1-morpholino-1-cyclohexene (3.88 ml, 22.6 mmol) was stirred at 70°C for 3 hours. The solvent was distilled out under reduced pressure. The resulting 20 residue was purified by alumina column chromatography (development solvent; ethyl acetate), to give 3-nitro-5,6,7,8-tetrahydroquinoline (2.42 g) as a powder from methanol - water (1:4).

¹H NMR (DMSO-d₆) δ: 1.87 (4H, m), 2.90 (4H, m), 8.15 (1H, s), 9.16 (1H, s).

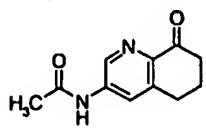
3) 10% Palladium-carbon (200 mg) was added to a methanol solution (68 ml) of 3-nitro-5,6,7,8-tetrahydroquinoline (2.41 g, 13.5 mmol) obtained in 2), which was stirred under hydrogen atmosphere for 16 hours. 30 After a catalyst was filtered off, the solvent was distilled out under reduced pressure. The resulting residue was dissolved in pyridine (35 ml). Anhydrous ethyl acetate (1.91 ml, 20.3 mmol) was added to the solution, which was stirred at room temperature for 1 hour. After 35 completion of the reaction, the solvent was distilled out

under reduced pressure. Diisopropyl ether - n-hexane (1:8) was added to the resulting residue, to give the titled compound (2.48 g) as a colorless powder.

¹H NMR (CDCl₃) δ : 1.80-1.87 (4H, m), 2.18 (3H, s), 2.77 (2H, m), 2.87 (2H, m), 7.72 (1H, br), 7.94 (1H, s), 8.24 (1H, s);

Reference Example 84

N-(8-Oxo-5,6,7,8-tetrahydro-3-quinolinyl)acetamide



10 1) m-Chloroperbenzoic acid (3.83 g, 15.5 mmol) was added to a chloroform solution (65 ml) of N-(5,6,7,8-tetrahydro-3-quinolinyl)acetamide (2.46 g, 12.9 mmol) obtained in Reference Example 83 under ice-cooling, which
15 was stirred at room temperature for 16 hours. After the solvent was distilled out under reduced pressure, the residue was powdered with ethyl acetate, to give N-(1-oxide-5,6,7,8-tetrahydro-3-quinolinyl)acetamide (2.00 g).

20 ¹H NMR (DMSO-d₆) δ : 1.64 (2H, m), 1.75 (2H, m), 2.04 (3H, s), 2.66 (4H, m), 7.13 (1H, s), 8.56 (1H, s), 10.12 (1H, s).

25 2) Anhydrous ethyl acetate (30 ml) was added to N-(1-oxide-5,6,7,8-tetrahydro-3-quinolinyl)acetamide (1.99 g, 9.65 mmol) obtained in 1), which was stirred at 80°C for 3 hours. The reaction mixture was cooled to room temperature. The solvent was distilled out under reduced pressure, and the resulting residue was purified by alumina column chromatography (development solvent; ethyl acetate). The resulting oily substance was dissolved in methanol (110 ml). 1N Sodium hydroxide (21.5 ml) was added to the solution under ice-cooling, which was stirred at room temperature for 1 hour. The solvent was distilled out under reduced pressure. Chloroform was added to the residue,

which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was purified by alumina column chromatography (development solvent; ethyl acetate:methanol = 5:1), to give N-(8-hydroxy-5,6,7,8-tetrahydro-3-quinolinyl)acetamide (1.08 g) as a powder from ethyl acetate and diisopropyl ether.

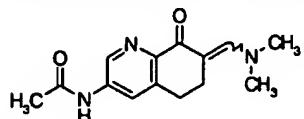
¹H NMR (CDCl₃) δ: 1.79 (2H, m), 1.96 (1H, m), 2.22 (3H, s), 2.24 (1H, m), 2.82 (2H, m), 4.69 (1H, m), 7.49 (1H, br), 7.92 (1H, s), 8.30 (1H, s).

3) Manganese dioxide (4.47 g, 51.4 mmol) was added to chloroform (26 ml) solution of N-(8-hydroxy-5,6,7,8-tetrahydro-3-quinolinyl)acetamide (1.06 g, 5.14 mmol) obtained in 2), which was stirred at room temperature for 1 day. After completion of the reaction, the insoluble matters were filtered off, and the filtrate was concentrated under reduced pressure. Diisopropyl ether and hexane were added to the resulting residue, to give the titled compound (858 mg) as a colorless powder.

¹H NMR (CDCl₃) δ: 2.20 (2H, m), 2.26 (3H, s), 2.77 (2H, m), 3.03 (2H, m), 8.10 (1H, br), 8.39 (1H, s), 8.42 (1H, s).

25 Reference Example 85

N-[7-[(Dimethylamino)methylidene]-8-oxo-5,6,7,8-tetrahydro-3-quinolinyl]acetamide

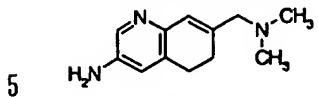


The titled compound was obtained by carrying out the same operation as in Reference Example 47, using N-(8-oxo-5,6,7,8-tetrahydro-3-quinolinyl)acetamide obtained in Reference Example 84.

¹H NMR (CDCl₃) δ: 2.09 (3H, s), 2.78 (2H, m), 2.85 (2H, m), 3.10 (6H, s), 7.55 (1H, s), 8.01 (1H, s), 8.56 (1H, s).

Reference Example 86

N-[(3-Amino-5,6-dihydro-7-quinolinyl)methyl]-N,N-dimethylamine

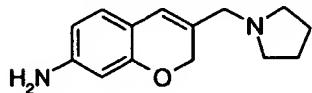


The titled compound was obtained by carrying out the same operation as in Reference Example 41-2), using N-[7-[(dimethylamino)methylidene]-8-oxo-5,6,7,8-tetrahydro-3-quinolinyl]acetamide obtained in Reference Example 85.

¹H NMR (CDCl₃) δ : 2.23 (6H, s), 2.33 (2H, t, J=8.1 Hz), 2.78 (2H, t, J=8.1 Hz), 2.99 (2H, s), 3.59 (2H, br), 6.43 (1H, s), 6.74 (1H, d, J=2.5 Hz), 7.84 (1H, d, J=2.5 Hz).

15 Reference Example 87

3-(1-Pyrrolidinylmethyl)-2H-chromen-7-amine



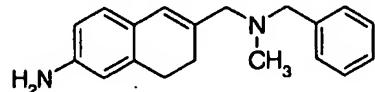
The titled compound was obtained as an oily substance by carrying out the same operations as in Example 41-1), Reference Example 52 and Example 41-2) in this order, using 7-acetylamino-3,4-dihydrochromen-4-one.

¹H-NMR (CDCl₃) δ : 1.77-179 (4H, m), 2.45-2.47 (4H, m), 3.11 (2H, s), 3.66 (2H, s), 4.74 (2H, s), 6.14-6.21 (3H, m), 6.75 (1H, d, J = 7.8 Hz).

25

Reference Example 88

6-[(N-Benzyl-N-methylamino)methyl]-7,8-dihydro-2-naphthalenamine



30 The titled compound was obtained as an oily substance by carrying out the same operation as in Reference Example 52, using 6-acetamido-2-(N,N-dimethylaminomethylidene)-

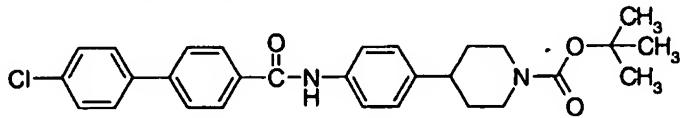
1-tetralone obtained in Example 41-1).

¹H-NMR (CDCl₃) δ : 2.17 (3H, s), 2.35 (2H, t, J = 8.1 Hz), 2.73 (2H, t, J = 8.1 Hz), 3.04 (2H, s), 3.48 (2H, s), 3.58 (2H, s), 6.29 (1H, s), 6.44 - 6.46 (2H, m), 6.82 (1H, d, J = 8.1 Hz), 7.03-7.45 (5H, m).

Reference Example 89

4'-Chloro-N-[4-(4-piperidinyl)phenyl][1,1'-biphenyl]-4-carboxamide

10



15

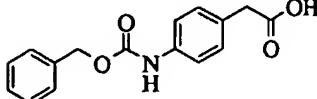
An ethanol solution (30 ml) of tert-butyl 4-(4-nitrophenyl)-1-piperidinecarboxylate (1.7 g) was subjected to catalytic hydrogenation using 10% palladium carbon (0.2 g) as a catalyst under normal temperature and normal pressure. After the catalyst was filtered off, the filtrate was concentrated to give tert-butyl 4-(4-aminophenyl)-1-piperidinecarboxylate as a viscous oily substance. The titled compound (2.2 g) was obtained as colorless crystals, by carrying out the same operation as in Example 1, using the resulting oily substance and

4'-chloro[1,1'-biphenyl]-4-carboxylic acid (1.43 g).
¹H-NMR (CDCl₃+ DMSO-d₆) δ : 1.05-1.32 (11H, m), 1.38-1.50 (2H, m), 2.20-2.50 (3H, m), 3.75-3.90 (2H, m), 6.81 (2H, d, J=8.4 Hz), 7.07 (2H, d, J=8.4 Hz), 7.20-7.36 (6H, m), 7.69 (2H, d, J=8.1Hz), 9.44 (1H, s).

Melting point: 232 - 233°C (crystallization solvent : ethyl acetate)

Reference Example 90

30 2-[4-[(Benzyl)carbonyl]amino]phenyl]ethyl acetate



To an ethyl acetate (100 ml) suspension of 4-

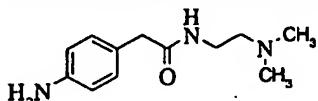
aminophenylethyl acetate (10 g), saturated aqueous sodium bicarbonate solution (100 ml) was added, and further, benzyloxycarbonyl chloride (12.3 ml) was added dropwise under ice-cooling. After stirring for 1 hour,

5 hydrochloric acid was added to the reaction mixture to make it acidic, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The residue was recrystallized from ethyl

10 acetate - hexane, to give the titled compound (17.3 g). Melting point: 148 - 149°C

Reference Example 91

2-(4-Aminophenyl)-N-[2-(dimethylamino)ethyl]acetamide

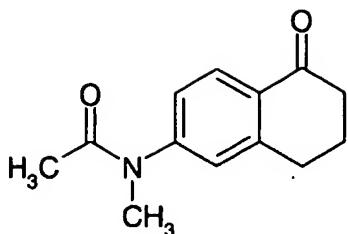


Pd-C (1 g) was added to a methanol (140 ml) solution of benzyl 4-[2-[2-(dimethylamino)ethyl]amino]-2-oxoethylphenylcarbamate (10 g), which was stirred under hydrogen atmosphere for 1 hour. Pd-C was removed, and the filtrate was concentrated. The residue was purified by alumina column chromatography (development solvent: ethyl acetate : hexane = 1:1), to give the titled compound (6.63 g) as an oily substance.

25 $^1\text{H-NMR}(\text{CDCl}_3)$ δ : 2.16 (6H, s), 2.05 (3H, s), 2.30-2.36 (2H, t, $J=6.2$ Hz), 3.23-3.32 (2H, dd, $J=11.4, 6.2$ Hz), 3.44 (2H, s), 6.00 (1H, s), 6.63-6.67 (2H, m), 7.00-7.07 (2H, m).

30 Reference Example 92

N-Methyl-N-(5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl)acetamide



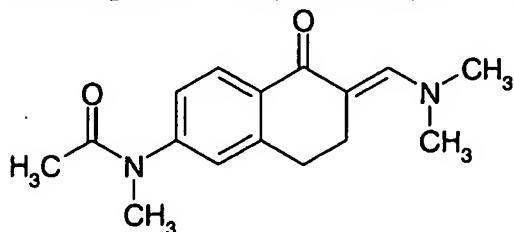
6-Acetamido-1-tetralone (10.0 g, 49.2 mmol) was dissolved in tetrahydrofuran (100 ml). Sodium hydride (oil, 3.0 g) was added to the solution, which was refluxed with heating under nitrogen atmosphere for 2 hours. After cooling, methyl iodide (30 ml) was added to the reaction mixture, which was refluxed with heating under nitrogen atmosphere for 2 hours. The reaction mixture was concentrated. Ethyl acetate and water were added to the residue, and extraction was conducted. The ethyl acetate layer was concentrated, and the residue was purified by alumina column chromatography (development solvent; ethyl acetate:n-hexane = 33:67 ~ 50:50). The product was concentrated under reduced pressure, and the residue was recrystallized from ethyl acetate - diisopropyl ether, to give the titled compound (4.3 g).

¹H-NMR (CDCl₃) δ: 1.96 (3H, brs), 2.18 (2H, m), 2.69 (2H, t, J=6.1 Hz), 2.99 (2H, t, J=5.9 Hz), 3.29 (3H, s), 7.01-7.15 (2H, m), 8.08 (1H, d, J=8.1 Hz).

20

Reference Example 93

N-[6-[(Dimethylamino)methylidene]-5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl]-N-methylacetamide



25

N-Methyl-N-(5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl)acetamide (4.3 g, 19.8 mmol) obtained in Reference Example 92 was dissolved in N,N-

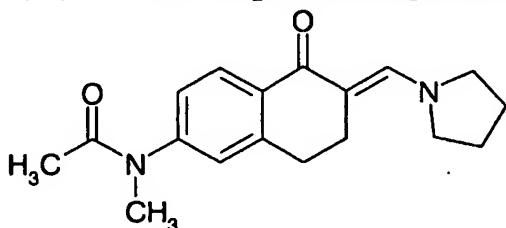
dimethylformamide dimethylacetal (50 ml), which was refluxed with heating under nitrogen atmosphere for 15 hours. The reaction mixture was concentrated, and the residue was washed with ethyl acetate and diisopropyl ether, to give the titled compound (3.9 g).

¹H-NMR (CDCl₃) δ : 1.93 (3H, brs), 2.84 (2H, dd, J=7.5, 5.6 Hz), 2.95 (2H, dd, J=7.5, 5.6 Hz), 3.16 (6H, s), 3.28 (3H, s), 6.99 (1H, s), 7.10 (1H, dd, J=8.1, 2.0 Hz), 7.75 (1H, s), 8.07 (1H, d, J=8.1 Hz).

10

Reference Example 94

N-Methyl-N-[5-oxo-6-[1-pyrrolidinylmethylidene]-5,6,7,8-tetrahydro-2-naphthalenyl]acetamide



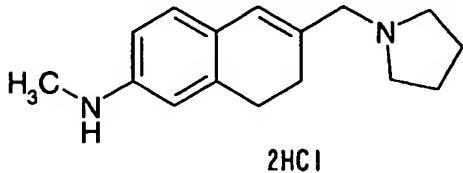
15 **N-[6-[(Dimethylamino)methylidene]-5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl]-N-methylacetamide** (5.7 g, 20.9 mmol) obtained in Reference Example 93 was dissolved in pyrrolidine (50 ml), which was refluxed with heating under nitrogen atmosphere for 3.5 hours. Then, ethyl acetate and water were added to the reaction mixture, and extraction was conducted. The ethyl acetate layer was concentrated, and the residue was recrystallized from ethyl acetate - diisopropyl ether, to give the titled compound (4.0 g, yield : 64%).

20 ¹H-NMR (CDCl₃) δ : 1.94 (7H, m), 2.84 (2H, dd, J=7.0, 5.6 Hz), 2.97 (2H, dd, J=7.0, 5.6 Hz), 3.28 (3H, s), 3.63 (4H, m), 6.98 (1H, s), 7.10 (1H, dd, J=8.1, 2.0 Hz), 7.95 (1H, s), 8.08 (1H, d, J=8.1 Hz).

25 **Reference Example 95**

N-Methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-

nephthalenamine dihydrochloride



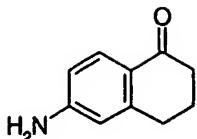
N-Methyl-N-[5-oxo-6-[1-pyrrolidinylmethylidene]-5,6,7,8-tetrahydro-2-naphthalenyl]acetamide (4.0 g, 13.4 mmol) obtained in Reference Example 94 was dissolved in methanol - ethyl acetate (10:1, 220 ml). 10% Palladium on carbon (50% wet, 0.4 g) was added to the solution, which was ice cooled. Stirring was begun under hydrogen atmosphere, and stirring was conducted for 2 days while returning the temperature of the reaction mixture to room temperature. A catalyst was filtered off, the reaction mixture was concentrated under reduced pressure, and the residue was dissolved in ethyl acetate. Extraction was conducted using 1N hydrochloric acid. The extract was made alkaline with 4N sodium hydroxide solution, and extraction was conducted using ethyl acetate. The extract was concentrated under reduced pressure. The residue was dissolved in tetrahydrofuran (100 ml) and 5N hydrochloric acid (100 ml), which was refluxed with heating for 13 hours.

The reaction mixture was concentrated. Ethyl acetate and saturated aqueous sodium carbonate solution were added to the residue, and extraction was conducted. The ethyl acetate layer was concentrated. 4N Hydrogen chloride - ethyl acetate solution was added to the resulting oily substance, which was concentrated. The residue was recrystallized from methanol - ethyl acetate, to give the titled compound (2.8 g, yield : 66%).

¹H-NMR (DMSO-d₆) δ: 1.98 (4H, m), 2.45 (4H, m), 2.81 (5H, m), 3.01 (2H, brd), 3.44 (2H, brd), 3.86 (2H, d, J=5.0 Hz), 7.02-7.10 (3H, m), 10.89 (1H, brs).

Reference Example 96

6-Amino-3,4-dihydro-1-(2H)-naphthalenone



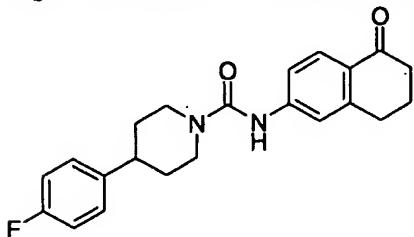
Concentrated hydrochloric acid (250 ml) was added to 6-acetamido-1-tetralone (20.0 g, 98.4 mmol), which was stirred at 100°C for 1 hour. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The residue was powdered with ethyl acetate and isopropyl ether, to give the titled compound (14.5 g).

^1H NMR (CDCl_3) δ : 2.07 (2H, m), 2.57 (2H, m), 2.83 (2H, m), 4.10 (2H, br), 6.42 (1H, d, $J=2.2$ Hz), 6.53 (1H, dd, $J=2.2$, 8.4 Hz), 7.89 (1H, d, $J=8.4$ Hz).

15

Reference Example 97

4-(4-Fluorophenyl)-N-(5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl)-1-piperidinecarboxamide

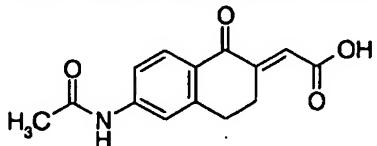


Pyridine (9.95 ml, 123 mmol) and 4-nitrophenyl chloroformate (12.4 g, 61.5 mmol) was added to a tetrahydrofuran (300 ml) solution of 6-amino-3,4-dihydro-1(2H)-naphthalenone (9.92 g, 61.5 mmol) obtained in Reference Example 96, which was stirred at room temperature for 3 hours. The solvent was distilled out under reduced pressure. 1N Hydrochloric acid was added to the residue to powder, which was washed with ethanol. 4N Aqueous sodium hydroxide solution was added to a dimethylsulfoxide (33 ml) solution of the resulting 4-nitrophenyl-5-oxo-

5,6,7,8-tetrahydro-2-naphthalenylcarbamate (2.20 g, 6.74 mmol) and 4-(4-fluorophenyl)piperidine hydrochloride (1.60 g, 7.42 mmol), which was stirred at room temperature for 1 hour. Ethyl acetate was added to the reaction 5 mixture, which was washed with 1N hydrochloric acid, aqueous potassium hydrogencarbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and the solvent was distilled out under reduced pressure. The resulting residue was purified by 10 alumina B column chromatography (development solvent; ethyl acetate), and powdered with isopropyl ether and hexane, to give the titled compound (1.89 g).

¹H NMR (CDCl₃) δ : 1.72 (2H, m), 1.92 (2H, m), 2.11 (2H, m), 2.61 (2H, m), 2.72 (1H, m), 2.93 (2H, m), 3.01 (2H, m), 4.23 15 (2H, m), 6.67 (1H, s), 7.00 (2H, m), 7.12 (3H, m), 7.61 (1H, s), 7.97 (1H, d, J=8.4 Hz).

Reference Example 98
[6-(Acetylamino)-1-oxo-3,4-dihydro-2(1H)- 20 naphthalenylidene]acetic acid

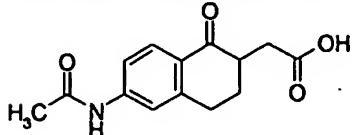


0.5N Aqueous sodium hydroxide solution (190 ml) was added to an aqueous solution (60 ml) of 6-acetamido-1-tetralone (5.00 g, 24.6 mmol) and glyoxylic acid (9.05 g, 98.5 mmol) under ice-cooling, which was stirred at 60°C for 25 16 hours. After cooling, concentrated hydrochloric acid was added to the reaction mixture. The precipitated crystals were collected, which was washed with water, to give the titled compound (3.73 g).

¹H NMR (DMSO-d₆) δ : 2.10 (3H, s), 2.95 (2H, m), 3.28 (2H, m), 6.63 (1H, s), 7.53 (1H, d, J=8.7Hz), 7.67 (1H, s), 7.91 30 (1H, d, J=8.7Hz), 10.32 (1H, s), 12.89 (1H, br).

Reference Example 99

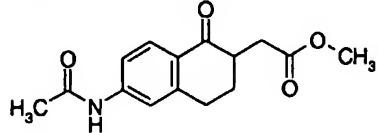
[6-(Acetylamino)-1-oxo-1,2,3,4-tetrahydro-2-naphthalenyl]acetic acid



5 70% Acetic acid - water solution (35 ml) of [6-(acetylamino)-1-oxo-3,4-dihydro-2(1H)-naphthalenyliden]acetic acid (3.50 g, 13.5 mmol) obtained in Reference Example 98 and zinc powder (2.1 g) was stirred at 100°C for 30 minutes. After cooling, zinc powder was
10 filtered. Ethyl acetate was added to the filtrate, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was purified by silica gel column
15 chromatography (development solvent; ethyl acetate : methanol = 10:1), and powdered with ethyl acetate and isopropyl ether, to give the titled compound (2.51 g).
1H NMR (CDCl₃) δ: 1.85-2.15 (2H, m), 2.08 (3H, s), 2.38 (1H, m), 2.71 (1H, m), 2.88 (2H, m), 3.05 (1H, m), 7.46 (1H, d, J=8.7Hz), 7.60 (1H, s), 7.80 (1H, d, J=8.7Hz), 10.21 (1H, s), 12.09 (1H, br).

Reference Example 100

25 Methyl [6-(acetylamino)-1-oxo-1,2,3,4-tetrahydro-2-naphthalenyl]acetate



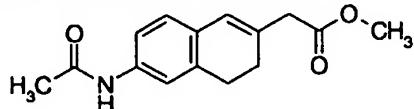
Methyl iodide (0.18 ml, 2.87 mmol) was added to a dimethylformamide solution (10 ml) of [6-(acetylamino)-1-oxo-1,2,3,4-tetrahydro-2-naphthalenyl]acetic acid (500 mg, 1.91 mmol) obtained in Reference Example 99 and potassium carbonate (529 mg, 3.82 mmol), which was stirred
30

at room temperature for 16 hours. Ethyl acetate was added to the reaction mixture, which was washed with aqueous sodium thiosulfate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and
5 then the solvent was distilled out under reduced pressure.

The resulting oily substance was purified by alumina B column chromatography (development solvent; ethyl acetate), to give the titled compound (527 mg).
¹H NMR (CDCl₃) δ: 1.98 (1H, m), 2.20 (3H, s), 2.23 (1H, m),
10 2.47 (1H, m), 3.30 (4H, m), 3.73 (3H, s), 7.21 (1H, d,
J=8.7Hz), 7.50-7.80 (2H, m), 7.97 (1H, d, J=8.7Hz).

Reference Example 101

Methyl [6-(acetylamino)-3,4-dihydro-2-
15 naphthalenyl]acetate



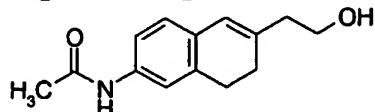
Sodium borohydride (72.4 mg, 1.91 mmol) was added to a methanol solution (10ml) of methyl [6-(acetylamino)-
1-oxo-1,2,3,4-tetrahydro-2-naphthalenyl]acetate (527 mg,
20 1.91 mmol) obtained in Reference Example 100 under ice-
cooling, which was stirred for 1 hour. Ethyl acetate was
added to the reaction mixture, which was washed with
saturated aqueous sodium chloride solution, dried over
anhydrous sodium sulfate, and then the solvent was
25 distilled out under reduced pressure. The resulting oily
substance was purified by alumina B column chromatography
(development solvent; ethyl acetate). Concentrated
sulfuric acid (0.14 ml) was added to an acetic acid solution
(7 ml) of the oil (404 mg, 1.46 mmol), which was stirred
30 at 40°C for 5 hours. The solvent was distilled out under
reduced pressure. Ethyl acetate was added to the residue,
which was washed with aqueous potassium carbonate solution
and saturated aqueous sodium chloride solution, dried over
anhydrous sodium sulfate, and then the solvent was

distilled out under reduced pressure. The resulting oily substance was purified by silica gel column chromatography (development solvent; hexane:ethyl acetate = 1:1), to give the titled compound (251 mg).

5 ¹H NMR (CDCl₃) δ : 2.16 (3H, s), 2.32 (2H, t, J=8.1Hz) , 2.82 (2H, t, J=8.1Hz), 3.21 (2H, s), 3.71 (3H, s), 6.30 (1H, s), 6.93 (1H, d, J=8.1Hz), 7.19 (2H, m) , 7.33 (1H, s).

Reference Example 102

10 N-[6-(2-Hydroxyethyl)-7,8-dihydro-2-naphthalenyl]acetamide

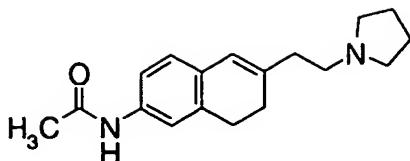


15 Lithium aluminum hydride (242 mg, 6.38 mmol) was added to a tetrahydrofuran solution (16 ml) of methyl [6-(acetylamino)-3,4-dihydro-2-naphthalenyl]acetate (827 mg, 3.19 mmol) obtained in Reference Example 101 under ice-cooling, which was stirred at room temperature for 1 hour. Ethyl acetate was added to the reaction mixture, which was washed with 1N hydrochloric acid and saturated 20 aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The residue was powdered with isopropyl ether, to give the titled compound (364 mg).

25 ¹H NMR (CDCl₃) δ : 1.43 (1H, m), 2.16 (3H, s), 2.26 (2H, t, J=8.1Hz) , 2.46 (2H, t, J=6.3Hz) , 2.81 (2H, t, J=8.1Hz), 3.78 (2H, m), 6.28 (1H, s), 6.94 (1H, d, J=8.1Hz), 7.08 (1H, br), 7.17 (1H, d, J=8.1Hz), 7.35 (1H, s).

Reference Example 103

30 N-[6-[2-(1-Pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl]acetamide

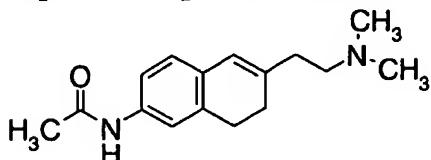


Methanesulfonyl chloride (0.131 ml, 1.69 mmol) was added to a dimethylformamide solution (7 ml) of N-[6-(2-hydroxyethyl)-7,8-dihydro-2-naphthalenyl]acetamide (355 mg, 1.53 mmol) obtained in Reference Example 102 and triethylamine (0.235 ml, 1.69 mmol) under ice-cooling, which was stirred for 30 minutes. Pyrrolidine (0.384 ml, 4.60 mmol) was added to the reaction mixture, which was stirred at 60 °C for 4 hours. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, and extraction was conducted using 1N hydrochloric acid. Potassium carbonate was added to the extract to make it alkaline, and extraction was conducted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and the solvent was distilled out under reduced pressure. The resulting residue was purified by alumina column chromatography (development solvent: ethyl acetate), to give the titled compound (294 mg).

¹H NMR (CDCl₃) δ : 1.79 (4H, m), 2.16 (3H, s), 2.25 (2H, m), 2.41 (2H, m), 2.55 (4H, m), 2.62 (2H, m), 2.78 (2H, m), 6.20 (1H, s), 6.91 (1H, d, J=8.1Hz), 7.18 (1H, d, J=7.8Hz), 7.32 (2H, m).

Reference Example 104

N-[6-(Dimethylamino)ethyl]-7,8-dihydro-2-naphthalenyl]acetamide



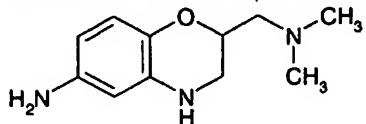
Methanesulfonyl chloride (0.0393 ml, 0.469 mmol) was added to a dimethylformamide solution (2 ml) of N-[6-

(2-hydroxyethyl)-7,8-dihydro-2-naphthalenyl]acetamide (102 mg, 0.426 mmol) obtained in Reference Example 102 and triethylamine (0.0652 ml, 0.469 mmol) under ice-cooling, which was stirred for 30 minutes. A tetrahydrofuran 5 solution (0.64 ml) of 2N dimethylamine was added to the reaction mixture, which was stirred at 60°C for 5 hours. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, and extraction was conducted using 1N hydrochloric acid. Potassium carbonate 10 was added to the extract to make it alkaline, and extraction was conducted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and the solvent was distilled out under reduced pressure. The resulting residue was purified by alumina column chromatography (development solvent; ethyl acetate), to give the titled compound (57.5 mg).

¹H NMR (CDCl₃) δ : 2.15 (3H, s), 2.24 (2H, m), 2.29 (6H, s), 2.36 (2H, m), 2.48 (2H, m), 2.78 (2H, m), 6.20 (1H, s), 6.90 20 (1H, d, J=8.1Hz), 7.20 (1H, d, J=8.1Hz), 7.35 (1H, s), 7.76 (1H, br).

Reference Example 105

6-Amino-2-[(dimethylamino)methyl]-1,4-benzoxazine



25 1) 2-Ethoxycarbonyl-6-nitro-1,4-benzoxazine (7.20 g, 0.029 mol) obtained by a known method by documents (Journal of heterocyclic chemistry, 19(5), p.1189 (1982)) was dissolved in methanol (50 ml). Sodium borohydride 30 (1.08 g, 0.029 mol) was added to the solution, which was stirred for 2 hours. The reaction mixture was concentrated. Ethyl acetate and aqueous potassium hydrogencarbonate solution were added to the residue, and extraction was conducted. The organic layer was washed

with water, and concentrated. A mixed solution of ethyl acetate and n-hexane (1:5) was added to the residue for crystallization. The crystallized product was collected by filtration, to give 2-hydroxymethyl-6-nitro-1,4-
5 benzoxazine (3.10 g) as a red powder.

¹H-NMR (CDCl₃) δ : 1.96 (1H, m), 3.34-3.49 (2H, m), 3.80-3.90 (2H, m), 4.09 (1H, brs), 4.30-4.40 (1H, m), 6.86 (1H, d, J=8.6 Hz), 7.50 (1H, d, J=2.8 Hz), 7.59 (1H, dd, J=2.8, 8.6 Hz).

10 2) 2-Hydroxymethyl-6-nitro-1,4-benzoxazine (1.00 g, 4.76 mmol) obtained in 1) and triethylamine (708 mg, 7.00 mmol) was dissolves in DMF (30 ml). Methanesulfonyl chloride (545 mg, 4.76 mmol) was added to the solution, which was stirred for 30 minutes. 50% Aqueous
15 dimethylamine solution (3 ml) was added to the reaction mixture, which was stirred at 70°C for 4 hours. Ethyl acetate and water were added to the mixture, and extraction was conducted. The organic layer was washed, and concentrated. The residue was subjected to alumina column
20 chromatography, and eluted with ethyl acetate:n-hexane (40:60), to give 2-[(dimethylamino)methyl]-6-nitro-1,4-benzoxazine (790 mg) as a colorless oily substance.

¹H-NMR (CDCl₃) δ : 2.33 (6H, s), 2.47-2.67 (2H, m), 3.19-3.25 (1H, m), 3.46-3.52 (1H, m), 4.09 (1H, brs), 4.30-4.35 (1H, m), 6.86 (1H, d, J=8.9 Hz), 7.48 (1H, d, J=2.8 Hz), 7.57 (1H, dd, J=2.8, 8.9 Hz).

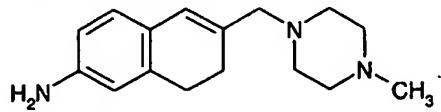
3) 2-[(Dimethylamino)methyl]-6-nitro-1,4-benzoxazine (760 mg, 3.2 mmol) obtained in 2) was dissolved in methanol (10 ml). Concentrated hydrochloric acid (3 ml) and iron powder (0.80 g) were added to the solution, which was stirred for 2 hours. The reaction mixture was concentrated. 1N Aqueous sodium hydroxide solution and ethyl acetate was added to the residue, and extraction was conducted. The organic layer was concentrated. The residue was subjected to alumina column chromatography, and eluted with ethyl acetate:n-hexane (20:80), to give the

titled compound (430 mg) as a colorless oily substance.
¹H-NMR (CDCl₃) δ : 2.31 (6H, s), 2.41-2.62 (2H, m), 3.12-3.17
(1H, m), 3.36-3.41 (1H, m), 3.30-3.50 (2H, brs), 3.67 (1H,
brs), 4.12-4.21 (1H, m), 5.99 (1H, d, J=2.5 Hz), 6.03 (1H,
5 dd, J=2.5, 8.4 Hz), 6.65 (1H, d, J=8.4 Hz).

Reference Example 106

6-[(4-Methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine

10



15

The titled compound was obtained by carrying out the same operation as in Reference Example 52, using 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone obtained in Example 41-1).

15

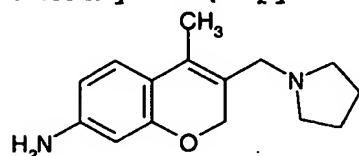
¹H NMR (CDCl₃) δ : 2.27 (2H, t, J=8.1 Hz), 2.29 (3H, s), 2.45 (8H, bs), 2.72 (2H, t, J=8.1 Hz), 3.03 (2H, s), 3.60 (2H, s), 6.26 (1H, s), 6.45-6.47 (2H, m), 6.80-6.83 (1H, m).

20

Reference Example 107

20

4-Methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine



25

The titled compound was obtained by carrying out the same operations as in Example 41-1) and Reference Example 69 in this order, using 1-acetylamo-3,4-dihydrochromen-1-one.

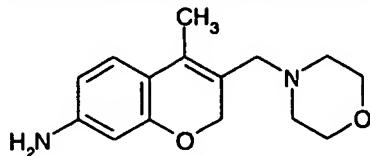
25

¹H NMR (CDCl₃) δ : 1.73-1.83 (4H, m), 1.99 (3H, s), 2.46-2.51 (4H, m), 3.22 (2H, s), 3.70 (2H, bs), 4.66 (2H, s), 6.18 (1H, d, J=2.2 Hz), 6.26 (1H, dd, J=2.2 Hz, 8.1 Hz), 7.00 (1H, d, J=8.1 Hz).

30

Reference Example 108

4-Methyl-3-(4-morpholinylmethyl)-2H-chromen-7-amine



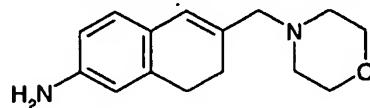
The titled compound was obtained by carrying out the same operations as in Example 41-1) and Reference Example 5 69 in this order, using 1-acetylamo-3,4-dihydrochromen-1-one.

^1H NMR (CDCl_3) δ : 1.98 (3H, s), 2.41-2.44 (4H, m), 3.08 (2H, s), 3.66-3.69 (6H, m), 4.62 (2H, s), 6.18 (1H, d, $J=2.2$ Hz), 6.26 (1H, dd, $J=2.2$ Hz, 8.1 Hz), 7.00 (1H, d, $J=8.1$ Hz).

10

Reference Example 109

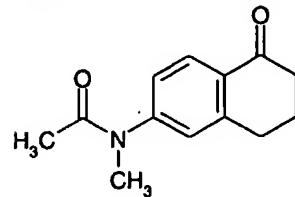
6-(4-Morpholinylmethyl)-7,8-dihydro-2-naphthalenamine



The titled compound was obtained by carrying out the 15 same operations as in Reference Example 52, using 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone obtained in Example 41-1).

$^1\text{H-NMR}$ (CDCl_3) δ : 2.28 (2H, t, $J=7.8$ Hz), 2.42 (4H, t, $J=4.4$ Hz), 2.72 (2H, t, $J=7.8$ Hz), 3.01 (2H, s), 3.60 (2H, brs.), 20 3.70 (4H, t, $J=4.4$ Hz), 6.26 (1H, s), 6.46 (2H, m), 6.82 (1H, d, $J=8.7$ Hz).

Reference Example 110
N-Methyl-N-(5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl)acetamide



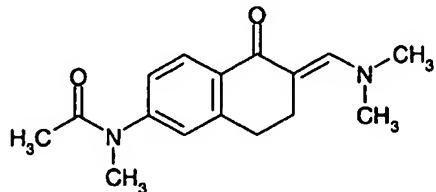
6-Acetamido-1-tetralone (13.7 g, 67.4 mmol) was dissolved in tetrahydrofuran (40 ml). Sodium

hydride(oil)(2.40 g, 101 mmol) was added to the solution, which was refluxed with heating for 2.5 hours. After cooling, methyl iodide(20 ml)was added to the reaction mixture, which was stirred at 40°C for 15 hours. The 5 reaction mixture was poured into a cold water, and extraction was conducted using ethyl acetate. The extract was washed with 1N hydrochloric acid and 1 N aqueous sodium hydroxide solution. The ethyl acetate layer was concentrated. The residue was purified by alumina column 10 chromatography (development solvent; ethyl acetate:n-hexane = 50:50 ~ 100:0) . The eluent was concentrated under reduced pressure. The resulting residue was recrystallized from ethyl acetate - diisopropyl ether, to give the titled compound(8.3 g).

15 $^1\text{H-NMR}$ (CDCl_3) δ : 1.96 (3H, s), 2.19(2H, m), 2.69 (2H, t, J=6.2 Hz), 2.99 (2H, t, J=5.9 Hz), 3.29 (3H, s), 7.10-7.15 (2H, m), 8.09 (1H, d, J=8.4 Hz).

Reference Example 111

20 $\text{N-[6-[(E)-(Dimethylamino)methylidene]-5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl]-N-methylacetamide}$



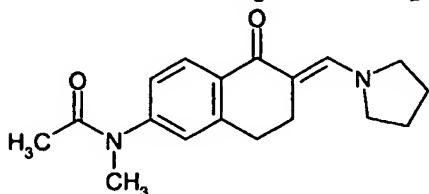
25 $\text{N-Methyl-N-(5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl)acetamide}$ (4.3 g, 19.8 mmol) obtained in Reference Example 110 was dissolved in $\text{N,N-dimethylformamide-dimethylacetal}$ (50 ml), which was refluxed with heating under nitrogen atmosphere for 15 hours. The reaction mixture was concentrated under reduced pressure. The resulting residue was washed with 30 ethyl acetate - diisopropyl ether, to give the titled compound(3.9 g).

$^1\text{H-NMR}$ (CDCl_3) δ : 1.93 (3H, s), 2.86 (2H, t, J=7.3 Hz), 2.95

(2H, t, J=7.3 Hz), 3.16 (6H, s), 3.28 (3H, s), 6.99 (1H, s), 7.09 (1H, d, J=8.1 Hz), 7.75 (1H, s), 8.07 (1H, d, J=8.1 Hz).

5 Reference Example 112

N-Methyl-N-[5-oxo-6-((E)-1-pyrrolidinylmethylidene)-5,6,7,8-tetrahydro-2-naphthalenyl]acetamide



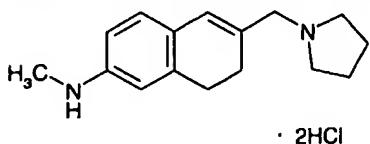
N-[6-[(E)-(Dimethylamino)methylidene]-5-oxo-

10 5,6,7,8-tetrahydro-2-naphthalenyl]-N-methylacetamide
(5.7 g, 20.9 mmol) obtained in Reference Example 111 was dissolved in pyrrolidine (50 ml), which was refluxed with heating under nitrogen atmosphere for 3.5 hours. The reaction mixture was poured into cold water, and extraction
15 was conducted using ethyl acetate. The ethyl acetate layer was concentrated. The resulting residue was recrystallized from ethyl acetate - diisopropyl ether, to give the titled compound (4.0 g).

¹H-NMR (CDCl₃) δ: 1.93-1.96 (7H, m), 2.85 (2H, t, J=6.7 Hz),
20 2.96 (2H, t, J=6.7 Hz), 3.28 (3H, s), 3.63 (4H, m), 6.99 (1H, s), 7.10 (1H, dd, J=8.4, 2.0 Hz), 7.95 (1H, s), 8.08 (1H, d, J=8.4 Hz).

Reference Example 113

25 N-Methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine dihydrochloride



· 2HCl

N-Methyl-N-[5-oxo-6-((E)-1-pyrrolidinylmethylidene)-5,6,7,8-tetrahydro-2-

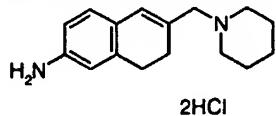
30 naphthalenyl]acetamide (4.0 g, 13.4 mmol) obtained in

Reference Example 112 was dissolved in methanol - acetic acid(10:1, 220 ml). 10% Palladium on carbon (0.4 g) was added to the solution, which was stirred under hydrogen atmosphere for 48 hours. The catalyst was filtered off, and 5 the reaction mixture was concentrated under reduced pressure. Ethyl acetate and 1N hydrochloric acid were added to the residue, and extraction was conducted. After the water layer was made alkaline with 4N aqueous sodium hydroxide solution, extraction was conducted using ethyl 10 acetate. The ethyl acetate layer was concentrated. Tetrahydrofuran - 5N hydrochloric acid (50:50, 200 ml) was added to the resulting residue, which was refluxed with heating for 13 hours. The reaction mixture was concentrated. Ethyl acetate and saturated aqueous sodium 15 carbonate solution was added to the residue, and extraction was conducted. 4N Hydrogen chloride - ethyl acetate solution was added to the ethyl acetate layer, which was concentrated under reduced pressure. The resulting residue was recrystallized from methanol - ethyl acetate, 20 to give the titled compound(2.8 g).

¹H-NMR (DMSO-d₆) δ : 1.98 (4H, m), 2.45 (4H, m), 2.81 (5H, m), 3.01 (2H, m), 3.44 (2H, m), 3.85 (1H, s), 3.86 (1H, s), 6.67 (1H, s), 7.02-7.10 (3H, m), 10.90 (1H, brs.).

25 Reference Example 114

6-(1-Piperidinylmethyl)-7,8-dihydro-2-naphthalenamine dihydrochloride



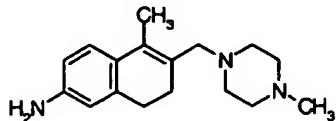
The titled compound was obtained by carrying out the 30 same operation as in Reference Example 52, using 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone obtained in Example 41-1).

¹H-NMR (DMSO-d₆) δ : 1.39 (1H, m), 1.80 (5H, m), 2.50 (5H, m), 2.83 (4H, m), 3.35-3.38 (2H, m), 3.79 (2H, s), 6.70 (1H,

s), 7.05-7.13 (3H, m), 10.40 (1H, brs).

Reference Example 115

5-Methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine

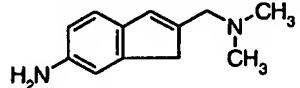


The titled compound was obtained by carrying out the same operation as in Reference Example 69, using 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone obtained in Example 41-1).

¹H NMR (CDCl₃) δ : 2.02 (3H, s), 2.27 (2H, t, J=8.1 Hz), 2.27 (3H, s), 2.44 (8H, bs), 2.63 (2H, t, J=8.1 Hz), 3.12 (2H, s), 3.61 (2H, s), 6.48-6.54 (2H, m), 7.08 (1H, d, J=7.8 Hz).

15 Reference Example 116

2-[(Dimethylamino)methyl]-1H-inden-6-amine



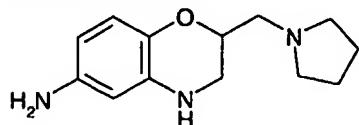
The titled compound was obtained by carrying out the same operation as in Example 41-2), using N-[2-[(E)-(dimethylamino)methylidene]-1-oxo-2,3-dihydro-1H-inden-5-yl]acetamide obtained in Reference Example 47.

¹H NMR (CDCl₃) δ : 2.24 (6H, s), 3.26 (2H, s), 3.33 (2H, s), ca.3.5 (2H, br), 6.58 (2H, m), 6.81 (1H, s), 7.08 (1H, d, J=8.1 Hz).

25

Reference Example 117

6-Amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine



30 A mixture of 6-nitro-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine and 4-(methylsulfonyl)-

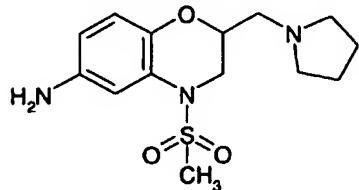
6-nitro-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine was obtained by carrying out the same operation as in Reference Example 105-2), using 2-hydroxymethyl-6-nitro-3,4-dihydro-2H-1,4-benzoxazine obtained in
5 Reference Example 105-1).

The titled compound was obtained by carrying out the same operation as in Reference Example 105-3), using the mixture obtained above.

¹H-NMR (CDCl₃) δ : 1.76-1.81 (4H, m), 2.50-2.70 (4H, m), 2.70
10 (2H, d, J=6.3Hz), 3.13-3.20 (1H, m), 3.20-3.40 (2H, brs),
3.39-3.43 (1H, m), 3.66 (1H, brs), 4.11-4.21 (1H, m), 5.99
(1H, d, J=2.7Hz), 6.03 (1H, dd, J=2.7, 8.4 Hz), 6.64 (1H,
d, J=8.4 Hz).

15 Reference Example 118

6-Amino-4-(methylsulfonyl)-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine

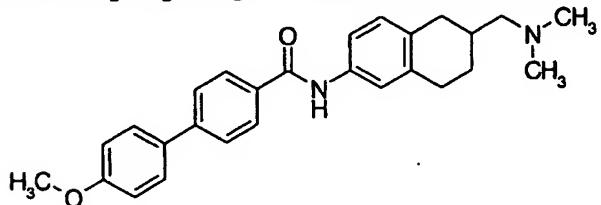


The titled compound was obtained by carrying out the
20 same operation as in Reference Example 105-3). using the
mixture of 6-nitro-2-(1-pyrrolidinylmethyl)-3,4-
dihydro-2H-1,4-benzoxazine and 4-(methylsulfonyl)-6-
nitro-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-
benzoxazine obtained in Reference Example 117.

¹H-NMR (CDCl₃) δ : 1.70-1.80 (4H, m), 2.50-2.70 (4H, m), 2.73
25 (2H, d, J=6.0Hz), 2.95 (3H, s), 3.21-3.29 (1H, m), 2.80-
3.10 (2H, brs), 4.10-4.21 (1H, m), 4.26-4.32 (1H, m), 6.43
(1H, dd, J=2.7, 8.4 Hz), 6.77 (1H, d, J=8.4 Hz), 7.11 (1H,
d, J=2.7Hz).

Example 1

N-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-(4'-methoxybiphenyl-4-yl)carboxamide



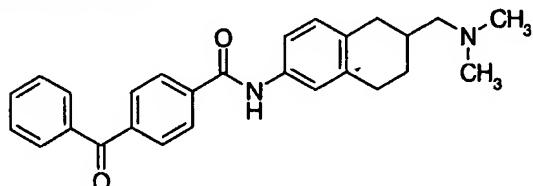
5 DMF solution (0.25 ml) of 2M HOBr, DMF solution (0.30 ml) of 2M WSCD, triethylamine (0.14 ml) and DMAP (0.132 g) were added to DMF solution (3 ml) of 6-amino-2-(N,N-dimethylamino)methyltetralin (0.139 g) and 4-(4-methoxyphenyl)benzoic acid (0.118 g). After the reaction mixture
10 was stirred at room temperature for 12 hours, 10% potassium carbonate solution was added, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting crude crystal
15 was washed with diethyl ether, which was recrystallized using ethyl acetate-hexane, to give the titled compound (0.124 g).

Melting point: 170 - 175°C.

20 Compounds described in the following Examples 2 and 3 were produced in the same manner as in Example 1.

Example 2

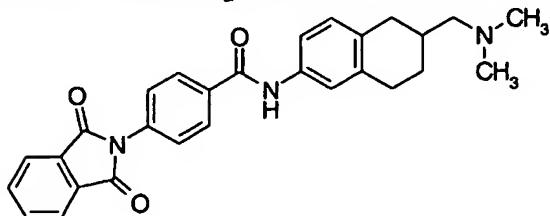
4-Benzoyl-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide



Melting point: 193 - 196°C (recrystallization solvent:
 ethyl acetate-hexane)

Example 3

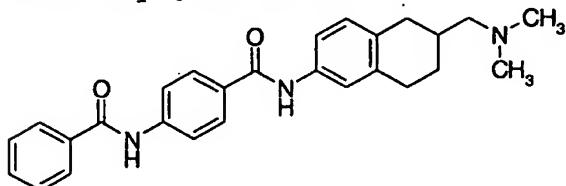
N-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-4-(1,3-dioxo-1,3-dihydro-2H-isoindol-2-yl) benzamide



5 Melting point: 235 - 240°C (washed with diethyl ether)

Example 4

4-(Benzoylamino)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide



10

6-Amino-2-(N,N-dimethylamino)methyltetralin hydrochloride (139 mg), 4-benzoylaminobenzoic acid (121 mg), WSCD (0.13 ml), HOBt (92 mg), triethylamine (0.14 ml) and DMAP (61 mg) were added to DMF (4 ml). After the reaction mixture was shaken at room temperature for 20 hours using a shaker, the reaction mixture was poured into water, and extraction was conducted using ethyl acetate-THF (1:1).

15 The organic layer was washed with water, saturated sodium bicarbonate solution and saturated aqueous sodium chloride solution, dried, and then concentrated. The resulting crude crystal was washed with hexane, to give the titled compound (181 mg).

20 Melting point : 241 - 242°C

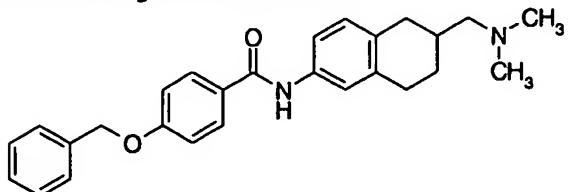
Washing solvent : hexane

25

Compounds described in the following Examples 5 to 14 were produced in the same manner as in Example 4.

Example 5

4-(Benzylxy)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide

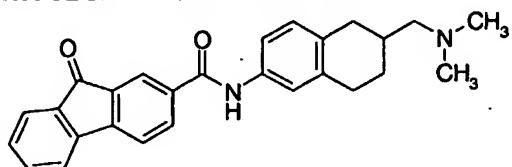


5 Melting point : 135 - 136°C

Washing solvent : hexane

Example 6

N-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-9-oxo-9H-fluoren-2-carboxamide

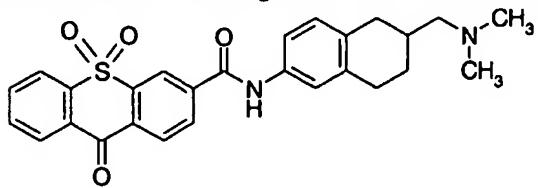


Melting point : 224 - 226°C

Washing solvent : hexane

Example 7

N-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-9,10,10-trioxo-9,10-dihydro-10¹-thioxanthene-3-carboxamide

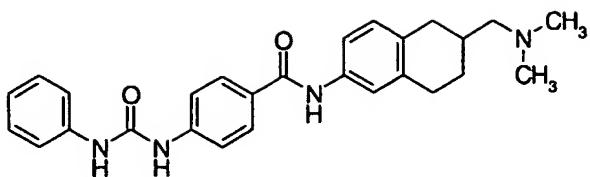


Melting point : 222 - 223°C (decomposition)

20 Washing solvent : hexane

Example 8

(4-Anilinocarbonyl)amino-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide

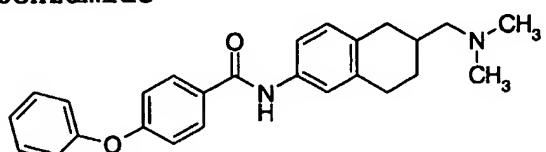


Melting point : 216 - 217°C (decomposition)

Washing solvent : hexane

5 Example 9

N-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-4-phenoxybenzamide

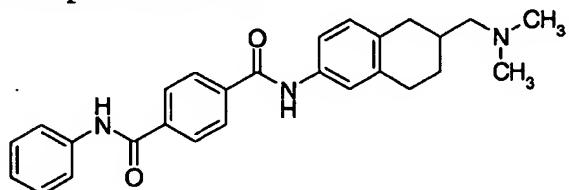


Melting point : 137 - 139°C

10 Washing solvent : hexane

Example 10

N¹-[2-(N,N-Dimethylamino)methyl-6-tetralinyl]-N⁴-phenylterephthalamide



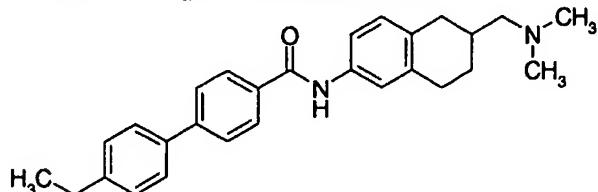
15

Melting point : 238 - 240°C (decomposition)

Washing solvent : hexane

Example 11

20 (4'-Ethylbiphenyl-4-yl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]carboxamide

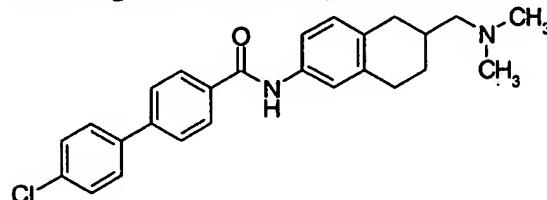


Melting point : 137 - 138°C

Washing solvent : hexane

Example 12

5 (4'-Chlorobiphenyl-4-yl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]carboxamide

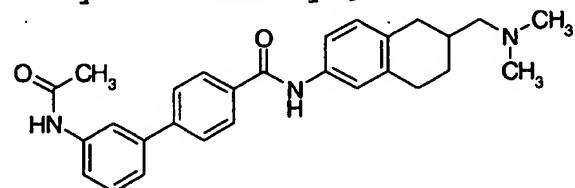


Melting point : 187 - 189°C

Washing solvent : hexane

10 **Example 13**

(4'-Acetylaminobiphenyl-4-yl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]carboxamide

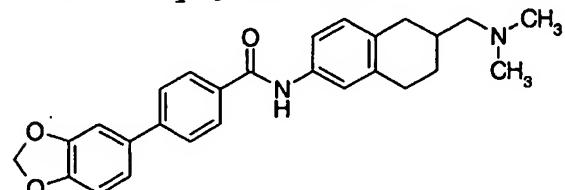


Melting point : 183 - 186°C

15 Washing solvent : hexane

Example 14

4-(1,3-Benzodioxol-5-yl)-N-[2-N,N-dimethylamino)methyl-6-tetralinyl]benzamide



20

Melting point : 174 - 176°C

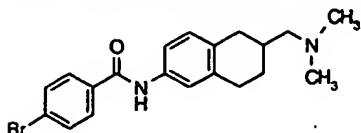
Washing solvent : hexane

Example 15

25 4-Bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-

tetrahydro-2-naphthalenyl]benzamide

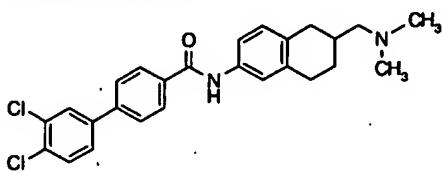
The titled compound was obtained as a white powder by the same method as in Example 1.



5 Melting point: 141 - 143°C (washing solvent: n-hexane)

Example 16

3',4'-Dichloro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



4-Bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (400 mg, 1.03 mmol) obtained in Example 15, 3,4-dichlorophenylboric acid (50 wt% THF-H₂O solution, 0.473 ml, 1.24 mmol), and 2N sodium carbonate solution (1.03 ml, 2.07 mmol) were dissolved in 50 ml of dimethoxyethane, then palladium tetrakis(triphenylphosphine) (35.8 mg, 0.031 mmol) was added under nitrogen atmosphere, which was stirred at 90°C for 15 hours.

Ethyl acetate was added to the reaction mixture, which was washed with saturated aqueous sodium chloride solution, dried using anhydrous magnesium sulfate, and the solvent was distilled out under reduced pressure. The residue was refined by alumina column chromatography (development solvent: n-hexane:ethyl acetate = 3:1), and pulverized with n-hexane to give the titled compound (204 mg) a white powder.

¹H-NMR (CDCl₃) δ: 1.41 (1H, m), 1.95 (2H, m), 2.26 (6H, s), 2.26-2.45 (3H, m), 2.83-2.99 (3H, m), 7.10 (1H, d, J=8.1 Hz), 7.26-7.77 (8H, m), 7.94 (2H, d, J=8.4 Hz).

Elemental analysis for $C_{26}H_{26}Cl_2N_2O \cdot 0.1H_2O$

Calcd. : C, 68.60; H, 5.80; N, 6.15.

Found: C, 68.42; H, 5.60; N, 5.92.

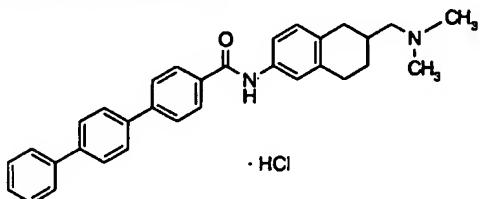
Melting point: 143 - 145°C (crystallization solvent:

5 ethyl acetate-hexane)

Example 17

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-phenyl[1,1'-biphenyl]-4-carboxamide

10 hydrochloride



The free basic substance (35 mg) of the titled compound was obtained in the same manner as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (400 mg, 1.03 mmol) obtained in Example 15, and 4-biphenylboric acid (1.25 g, 1.25 mmol). The resulting free basic substance (30 mg) was dissolved in 10 ml of methanol, then 100 ml of 1N hydrochloric acid was added, and the reaction mixture was stirred. The reaction mixture was concentrated, and pulverized using diethyl ether, to give the titled compound (35.3 mg) as a white powder.

¹H-NMR (DMSO-d₆, free base) δ: 1.32 (1H, m), 1.93 (2H, m), 2.15 (6H, s), 2.15-2.36 (3H, m), 2.74-2.94 (3H, m), 7.05 (1H, d, J=8.4 Hz), 7.40-7.55 (5H, m), 7.73-7.91 (8H, m), 8.07 (2H, d, J=8.4 Hz), 10.14 (1H, s).

Elemental analysis for $C_{32}H_{32}N_2O \cdot HCl \cdot 2H_2O$

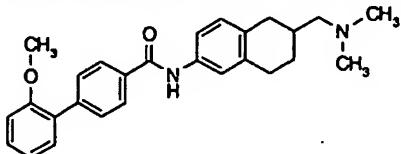
Calcd. : C, 72.10; H, 7.00; N, 5.25.

Found: C, 71.81; H, 6.57; N, 5.08.

30 Melting point: 220°C (decomposition) (crystallization solvent: methanol-diethyl ether)

Example 18

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-2'-methoxy[1,1'-biphenyl]-4-carboxamide



5 The titled compound (208 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) obtained in Example 15, and 2-methoxyphenylboric acid (118 mg, 0.775 mmol).

¹H-NMR (CDCl₃) δ : 1.42 (1H, m), 1.96 (2H, m), 2.23 (6H, s), 2.23-2.47 (3H, m), 2.85 (3H, m), 3.83 (3H, s), 7.05 (3H, m), 7.34 (3H, m), 7.47 (1H, s), 7.64 (2H, d, J=8.4 Hz), 7.79 (1H, s), 7.90 (2H, d, J=8.4 Hz).

15 Elemental analysis for C₂₇H₃₀N₂O₂ · 0.1H₂O

Calcd.: C, 77.89; H, 7.31; N, 6.73.

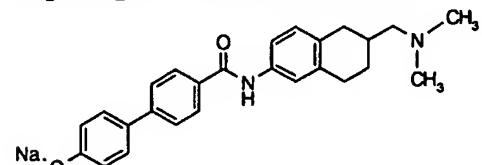
Found: C, 77.86; H, 7.18; N, 6.79.

Melting point: 155 - 157°C (crystallization solvent:
ethyl acetate-hexane)

20

Example 19

Sodium salt of N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-oxy[1,1'-biphenyl]-4-carboxamide



25

The titled compound (117 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) and 4-hydroxyphenylboric acid (107 mg, 0.775 mmol).

¹H-NMR (DMSO-d₆) δ : 1.36 (1H, m), 1.89 (2H, m), 2.15 (6H, s), 2.15-2.35 (3H, m), 2.77 (3H, m), 6.88 (2H, d, J=8.4 Hz), 7.02 (1H, d, J=8.4 Hz), 7.48 (1H, d, J=8.4 Hz), 7.53 (1H, s), 7.59 (2H, d, J=8.4 Hz), 7.73 (2H, d, J=8.4 Hz), 8.00 (2H, d, J=8.4 Hz), 10.07 (1H, s).

Elemental analysis for C₂₆H₂₇N₂O₂Na · 0.2H₂O

Calcd. : C, 73.29; H, 6.48; N, 6.59.

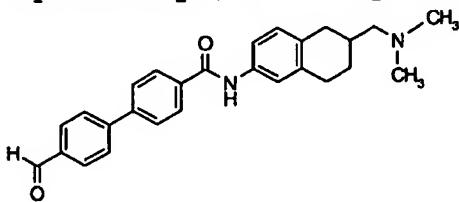
Found : C, 73.25; H, 6.18; N, 6.36.

Melting point: 246 - 248°C (crystallization solvent:

10 ethyl acetate-diethyl ether)

Example 20

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-formyl[1,1'-biphenyl]-4-carboxamide



15

The titled compound (205 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) and 4-formylphenylboric acid (145 mg, 0.968 mmol).

20 ¹H-NMR (CDCl₃) δ : 1.41 (1H, m), 1.95 (2H, m), 2.26 (6H, s), 2.26-2.42 (3H, m), 2.85-2.94 (3H, m), 7.09 (2H, d, J=8.1 Hz), 7.32 (1H, d, J=8.4 Hz), 7.47 (1H, m), 7.63-7.94 (3H, m), 7.87-7.99 (4H, m), 8.13 (1H, s), 10.11 (1H, s).

25 Elemental analysis for C₂₇H₂₈N₂O₂ · 0.2H₂O

Calcd. : C, 77.93; H, 6.88; N, 6.73.

Found : C, 77.89; H, 6.75; N, 6.71.

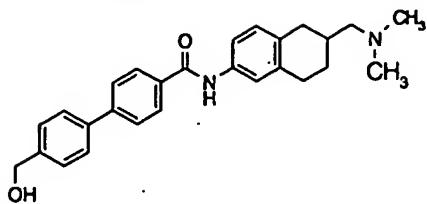
Melting point: 130 - 132°C (crystallization solvent: ethyl acetate-diethyl ether)

30

Example 21

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-

naphthalenyl]-4'-(hydroxymethyl)[1,1'-biphenyl]-4-carboxamide



5 N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-formyl[1,1'-biphenyl]-4-carboxamide (100 mg, 0.242 mmol) was dissolved in tetrahydrofuran-methanol (1:1) solution (2.4 ml), then sodium borohydride (18.3 mg, 0.485 mmol) was added, which was stirred for 2 hours. Ethyl acetate was added to the
 10 reaction mixture, which was washed with saturated aqueous sodium chloride solution, dried using anhydrous magnesium sulfate, and the solvent was distilled out under reduced pressure. The residue was pulverized using ether-n-hexane, to give the titled compound (86 mg) as a white
 15 powder.

¹H-NMR (CDCl₃) δ : 1.39 (1H, m), 1.94 (2H, m), 2.25 (6H, s), 2.25-2.44 (3H, m), 2.82-2.95 (3H, m), 4.78 (2H, s), 7.07 (1H, d, J=8.4 Hz), 7.31 (1H, d, J=8.4 Hz), 7.38-7.56 (4H, m), 7.64-7.70 (3H, m), 7.85 (1H, s), 7.93 (2H, d, J=8.4 Hz).

20 Elemental analysis for C₂₇H₃₀N₂O₂ · 0.2H₂O

Calcd. : C, 77.56; H, 7.33; N, 6.70.

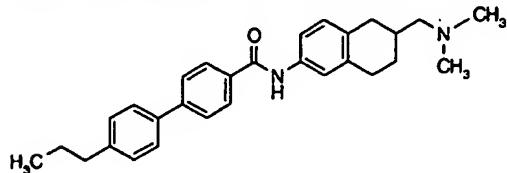
Found : C, 77.53; H, 7.27; N, 6.55.

Melting point: 138 - 139°C (crystallization solvent: ethyl acetate-diethyl ether)

25

Example 22

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-propyl[1,1'-biphenyl]-4-carboxamide



The titled compound (158 mg) was obtained as a white powder by the same method as in Example 1, using N-[(6-amino-1,2,3,4-tetrahydro-2-naphthalenyl)methyl]-N,N-dimethylamine (102 mg, 0.499 mmol), and 4-(4-propyl)benzoic acid (144 mg, 0.599 mmol).

¹H-NMR (CDCl₃) δ : 0.98 (3H, t, J=7.5 Hz), 1.40 (1H, m), 1.69 (2H, m), 1.94 (2H, m), 2.25 (6H, s), 2.25-2.45 (3H, m), 2.64 (2H, t, J=7.5 Hz), 2.85 (3H, m), 7.08 (1H, d, J=7.8 Hz), 7.26 (3H, m), 7.46 (1H, s), 7.54 (2H, d, J=8.1 Hz), 7.67 (2H, d, J=8.1 Hz), 7.81 (1H, s), 7.91 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₉H₃₄N₂O

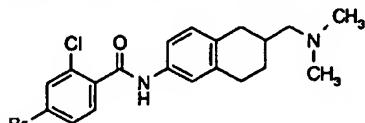
Calcd.: C, 81.65; H, 8.03; N, 6.57.

Found: C, 81.30; H, 7.94; N, 6.40.

Melting point: 186 - 188°C (crystallization solvent: ethyl acetate-diethyl ether)

Example 23

4-Bromo-2-chloro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide



The titled compound (483 mg) was obtained as a white powder by the same method as in Example 1, using N-[(6-amino-1,2,3,4-tetrahydro-2-naphthalenyl)methyl]-N,N-dimethylamine (300 mg, 1.47 mmol) and 4-bromo-2-chloro benzoic acid (415 mg, 1.76 mmol).

¹H-NMR (CDCl₃) δ : 1.40 (1H, m), 1.94 (2H, m), 2.25 (6H, s), 2.25-2.44 (3H, m), 2.94 (3H, m), 7.08 (1H, d, J=8.4 Hz), 7.28 (1H, m), 7.41 (1H, s), 7.50 (1H, m), 7.61 (2H, m), 7.81 (1H, s).

Elemental analysis for C₂₀H₂₂BrClN₂O

Calcd.: C, 56.96; H, 5.26; N, 6.64.

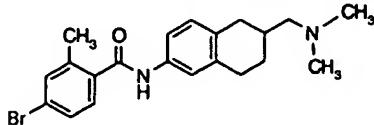
Found: C, 57.09; H, 5.37; N, 6.55.

Melting point: 130 - 132°C (crystallization solvent: ethyl acetate-diethyl ether)

Example 24

4-Bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-2-methylbenzamide

5



10

The titled compound (418 mg) was obtained as a white powder by the same method as in Example 1, using N-[(6-amino-1,2,3,4-tetrahydro-2-naphthalenyl)methyl]-N,N-dimethylamine (293 mg, 1.43 mmol) and 4-bromo-2-methyl

10
benzoic acid (370 mg, 1.72 mmol).

¹H-NMR (CDCl₃) δ : 1.40 (1H, m), 2.04 (2H, m), 2.25 (6H, s), 2.25-2.40 (3H, m), 2.46 (3H, s), 2.88 (3H, m), 7.07 (1H, d, J=7.8 Hz), 7.21-7.41 (6H, m).

Elemental analysis for C₂₁H₂₅BrN₂O

15

Calcd. : C, 62.85; H, 6.28; N, 6.98.

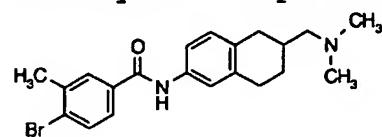
Found : C, 63.10; H, 6.11; N, 6.97.

Melting point: 140 - 142°C (crystallization solvent: ethyl acetate-hexane)

20

Example 25

4-Bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-3-methylbenzamide



25

The titled compound (434 mg) was obtained as a white powder by the same method as in Example 1, using N-[(6-amino-1,2,3,4-tetrahydro-2-naphthalenyl)methyl]-N,N-dimethylamine (300 mg, 1.47 mmol) and 4-bromo-3-methyl benzoic acid (379 mg, 1.76 mmol).

30

¹H-NMR (CDCl₃) δ : 1.40 (1H, m), 1.93 (2H, m), 2.25 (6H, s), 2.25-2.40 (3H, m), 2.46 (3H, s), 2.87 (3H, m), 7.07 (1H, d, J=7.8 Hz), 7.29 (1H, m), 7.40 (1H, s), 7.49 (1H, m), 7.61 (1H, d, J=8.1 Hz), 7.72 (2H, s-like).

Elemental analysis for $C_{21}H_{25}BrN_2O$

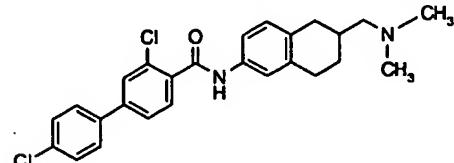
Calcd.: C, 62.85; H, 6.28; N, 6.98.

Found: C, 62.84; H, 6.05; N, 6.93.

Melting point: 154 - 155°C (crystallization solvent: ethyl
5 acetate-hexane)

Example 26

3,4'-Dichloro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



10

The titled compound (122 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-2-chloro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg,
15 0.607 mmol) obtained in Example 23, and 4-chlorophenyl boric acid (114 mg, 0.729 mmol).

1H -NMR ($CDCl_3$) δ : 1.41 (1H, m), 1.95 (2H, m), 2.26 (6H, s),
2.26-2.42 (3H, m), 2.85 (3H, m), 7.10 (1H, d, $J=8.4$ Hz),
7.31 (1H, m), 7.43-7.63 (8H, m), 7.87 (1H, d, $J=8.1$ Hz).

20 Elemental analysis for $C_{26}H_{26}Cl_2N_2O$

Calcd.: C, 68.87; H, 5.78; N, 6.18.

Found: C, 68.61; H, 5.49; N, 6.10.

Melting point: 177 - 179°C (crystallization solvent: ethyl acetate-diethyl ether)

25

Example 27

4'-Chloro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-3-methyl[1,1'-biphenyl]-4-carboxamide



30

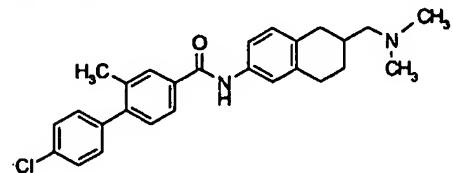
The titled compound (129 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-2-methylbenzamide (250 mg, 5 0.623 mmol) obtained in Example 24, and 4-chlorophenylboric acid (117 mg, 0.747 mmol).
¹H-NMR (CDCl₃) δ : 1.42 (1H, m), 1.96 (2H, m), 2.37 (6H, s), 2.37-2.47 (3H, m), 2.56 (3H, s), 2.90 (3H, m), 7.08 (1H, d, J=8.1 Hz), 7.26 (1H, m), 7.41 (6H, m), 7.53 (3H, m).
10 Elemental analysis for C₂₇H₂₉ClN₂O · H₂O
Calcd. : C, 71.90; H, 6.93; N, 6.21.
Found : C, 71.92; H, 6.52; N, 5.92.
Melting point: 163 - 165°C (crystallization solvent: ethyl acetate-diethyl ether)

15

Example 28

4'-Chloro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-2-methyl[1,1'-biphenyl]-4-carboxamide

20

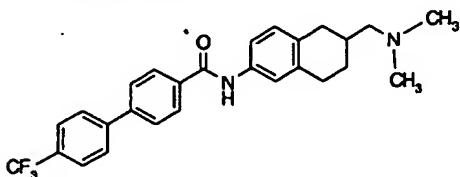


The titled compound (168 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-3-methylbenzamide (250 mg, 25 0.623 mmol) obtained in Example 25, and 4-chlorophenylboric acid (117 mg, 0.747 mmol).
¹H-NMR (CDCl₃) δ : 1.41 (1H, m), 1.95 (2H, m), 2.26 (6H, s), 2.24-2.42 (3H, m), 2.33 (3H, s), 2.85 (3H, m), 7.09 (1H, d, J=8.4 Hz), 7.26 (4H, m), 7.43 (3H, m), 7.73 (3H, m).
30 Elemental analysis for C₂₇H₂₉ClN₂O · 0.2H₂O
Calcd. : C, 74.28; H, 6.79; N, 6.42.
Found : C, 74.27; H, 6.73; N, 6.27.
Melting point: 193 - 195°C (crystallization solvent: ethyl

acetate-diethyl ether)

Example 29

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-(trifluoromethyl)[1,1-biphenyl]-4-carboxamide



The titled compound (194 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) obtained in Example 15, and 4-trifluoromethylphenylboric acid (147 mg, 0.775 mmol).

¹H-NMR (CDCl₃) δ: 1.41 (1H, m), 1.95 (2H, m), 2.25 (6H, s), 2.25-2.45 (3H, m), 2.89 (3H, m), 7.09 (1H, d, J=8.1 Hz), 7.31 (1H, d, J=8.1 Hz), 7.46 (1H, s), 7.70 (6H, m), 7.80 (1H, m), 7.96 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₇H₂₇F₃N₂O

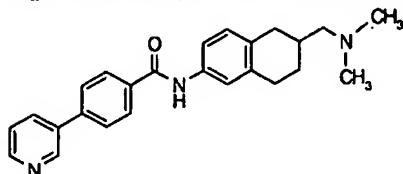
Calcd.: C, 71.66; H, 6.01; N, 6.19.

Found: C, 71.44; H, 6.05; N, 6.09.

Melting point: 205 - 206°C (crystallization solvent: ethyl acetate-diisopropyl ether)

Example 30

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4-(3-pyridinyl)benzamide



The titled compound (194 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-

tetrahydro-2-naphthalenyl)benzamide (250 mg, 0.645 mmol) obtained in Example 15, and 2-(3-pyridyl)-1,3,2,-dioxaborinane (126 mg, 0.775 mmol).

¹H-NMR (CDCl₃) δ : 1.41 (1H, m), 1.95 (2H, m), 2.26 (6H, s), 5 2.26-2.42 (3H, m), 2.85 (3H, m), 7.09 (1H, d, J=7.8 Hz), 7.30-7.47 (3H, m), 7.69 (2H, d, J=8.4 Hz), 7.86-7.99 (4H, m), 8.64 (1H, m), 8.87 (1H, m).

Elemental analysis for C₂₅H₂₇N₃O · 0.1H₂O

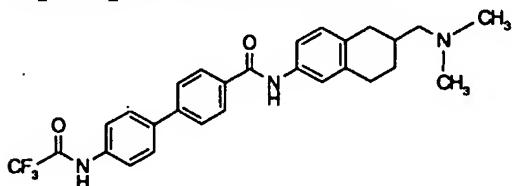
Calcd. : C, 77.53; H, 7.08; N, 10.85.

10 Found : C, 77.42; H, 7.05; N, 10.58.

Melting point: 177 - 178°C (crystallization solvent: ethyl acetate-diisopropyl ether)

Example 31

15 N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-[(trifluoroacetyl)amino][1,1'-biphenyl]-4-carboxamide



The titled compound (1.02 g) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (1.00 g, 2.58 mmol) obtained in Example 15, and 4-trifluoroacetamidophenylboric acid (722 mg, 3.10 mmol).

25 ¹H-NMR (CDCl₃) δ : 1.41 (1H, m), 2.05 (2H, m), 2.26 (6H, s), 2.26-2.42 (3H, m), 2.89 (3H, m), 7.09 (1H, d, J=8.4 Hz), 7.29 (2H, m), 7.46 (1H, s), 7.69 (7H, m), 7.94 (2H, d, J=8.1 Hz).

Elemental analysis for C₂₈H₂₈F₃N₃O₂

30 Calcd. : C, 67.87; H, 5.70; N, 8.48.

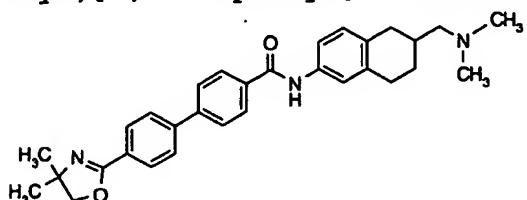
Found : C, 67.70; H, 5.53; N, 8.42.

Melting point: 235 - 237°C (crystallization solvent: ethyl

acetate-diisopropyl ether)

Example 32

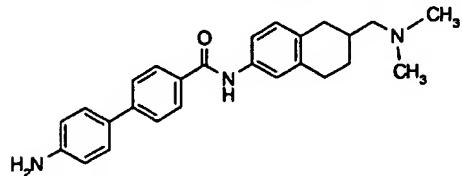
5 N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-(4,4-dimethyl-4,5-dihydro-1,3-oxazol-2-yl)[1,1'-biphenyl]-4-carboxamide



The titled compound (238 mg) was obtained as a white powder by the same method as in Example 16, using 4-
10. bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) obtained in Example 15, and 4-(4,4-dimethyl-4,5-dihydro-1,3-oxazol-2-yl)phenylboronic acid (170 mg, 0.775 mmol).
15. $^1\text{H-NMR}$ (CDCl_3) δ : 1.41 (7H, m), 1.94 (2H, m), 2.25 (6H, s), 2.25-2.41 (3H, m), 2.84 (3H, m), 4.14 (2H, s), 7.08 (1H, d, $J=7.8$ Hz), 7.30 (1H, m), 7.46 (1H, s), 7.68 (5H, m), 7.94 (2H, d, $J=8.4$ Hz), 8.03 (2H, d, $J=8.4$ Hz).
Elemental analysis for $\text{C}_{31}\text{H}_{35}\text{N}_3\text{O}_2 \cdot 0.2\text{H}_2\text{O}$
20. Calcd. : C, 76.74; H, 7.35; N, 8.66.
Found : C, 76.70; H, 7.19; N, 8.49.
Melting point: 185 - 187°C (crystallization solvent: ethyl acetate-diisopropyl ether)

25 **Example 33**

4'-Amino-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-

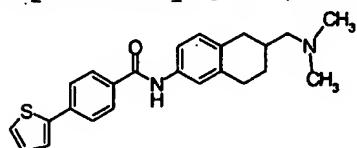
tetrahydro-2-naphthalenyl]-4'-[(trifluoroacetyl)amino][1,1'-biphenyl]-4-carboxamide (850 mg, 1.72 mmol) obtained in Example 31 was suspended in a mixed solution of methanol (8 ml) and tetrahydrofuran 5 (4 ml), then 1N sodium hydroxide (3.4 ml) was added, which was stirred at 50°C for 16 hours. The solvent was distilled out under reduced pressure, and the residue was pulverized using water, to give the titled compound (685 mg) as a white powder.

10 $^1\text{H-NMR}$ (CDCl_3) δ : 1.31 (1H, m), 1.89 (2H, m), 2.15 (6H, s), 2.15-2.34 (3H, m), 2.83 (3H, m), 5.36 (2H, s), 6.67 (2H, d, $J=8.4$ Hz), 7.03 (1H, d, $J=8.1$ Hz), 7.48 (4H, m), 7.68 (2H, d, $J=8.1$ Hz), 7.96 (2H, d, $J=8.4$ Hz), 10.02 (1H, s). Elemental analysis for $\text{C}_{26}\text{H}_{29}\text{N}_3\text{O} \cdot 1.1\text{H}_2\text{O}$

15 Calcd. : C, 74.47; H, 7.50; N, 10.02.
Found : C, 74.39; H, 7.41; N, 9.82.
Melting point: 148 - 150°C (crystallization solvent:
methanol-water)

20 Example 34

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4-(2-thienyl) benzamide



The titled compound (70 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) obtained in Example 15, and 2-thienylboric acid (99.1 mg, 0.775 mmol).

30 $^1\text{H-NMR}$ (CDCl_3) δ : 1.41 (1H, m), 1.94 (2H, m), 2.25 (6H, s), 2.25-2.45 (3H, m), 2.89 (3H, m), 7.11 (2H, m), 7.29-7.45 (4H, m), 7.71 (3H, m), 7.87 (2H, d, $J=8.4$ Hz).
Elemental analysis for $\text{C}_{24}\text{H}_{26}\text{N}_2\text{OS}$

Calcd.: C, 73.81; H, 6.71; N, 7.17.

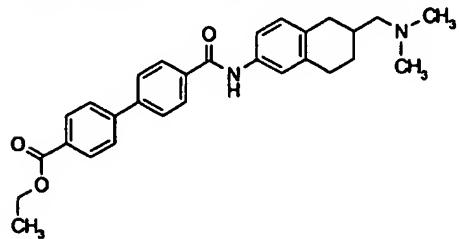
Found: C, 73.49; H, 6.59; N, 7.14.

Melting point: 165 - 166°C (crystallization solvent: ethyl acetate-diisopropyl ether)

5

Example 35

Ethyl 4'-[[[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]amino]carbonyl][1,1'-biphenyl]-4-carboxylate



10

The titled compound (202 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) obtained in Example 15, and 4-ethoxycarbonylphenylboric acid (150 mg, 0.775 mmol).

¹H-NMR (CDCl₃) δ: 1.42 (4H, m), 1.95 (2H, m), 2.26 (6H, s), 2.26-2.42 (3H, m), 2.89 (3H, m), 4.41 (2H, q, J=7.2 Hz), 7.09 (1H, d, J=8.4 Hz), 7.31 (1H, d, J=8.4 Hz), 7.47 (1H, s), 7.70 (4H, m), 7.80 (1H, s), 7.96 (2H, d, J=8.4 Hz), 8.14 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₉H₃₂N₂O₃

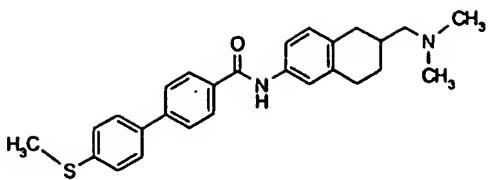
Calcd.: C, 76.29; H, 7.06; N, 6.14.

Found: C, 76.25; H, 7.07; N, 6.09.

Melting point: 156 - 158°C (crystallization solvent: ethyl acetate-diisopropyl ether)

Example 36

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-(methylsulfanyl)[1,1'-biphenyl]-4-carboxamide



The titled compound (360 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (500 mg, 1.29 mmol) obtained in Example 15, and 4-methylthiophenylboric acid (260 mg, 1.55 mmol).

¹H-NMR (CDCl₃) δ : 1.41 (1H, m), 1.94 (2H, m), 2.26 (6H, s), 2.26-2.42 (3H, m), 2.53 (3H, s), 2.94 (3H, m), 7.09 (1H, d, J=8.1 Hz), 7.29-7.36 (3H, m), 7.46 (1H, s), 7.56 (2H, d, J=8.4 Hz), 7.67 (2H, d, J=8.1 Hz), 7.78 (1H, m), 7.92 (2H, d, J=9.0 Hz).

Elemental analysis for C₂₇H₃₀N₂OS · 0.2H₂O

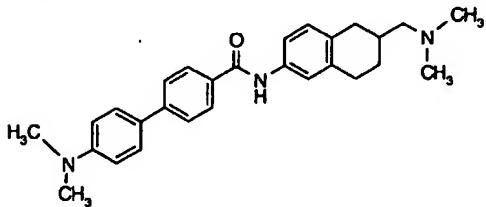
Calcd.: C, 74.69; H, 7.04; N, 6.45.

15 Found: C, 74.63; H, 7.03; N, 6.11.

Melting point: 178 - 180°C (crystallization solvent: ethyl acetate-diisopropyl ether)

Example 37

20 4'-(N,N-Dimethylamino)-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



25 4'-Amino-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide (150 mg, 0.375 mmol) obtained in Example 33, and paraformaldehyde (45.1 mg, 1.50 mmol) were suspended in mixed solution of methanol (1 ml) and tetrahydrofuran (1 ml). Sodium cyanohydroborate (94.4 mg, 1.50 mmol) was

added to the reaction mixture, which was stirred at 40°C for 18 hours. Ethyl acetate was added to the reaction mixture, which was washed with saturated aqueous sodium chloride solution, dried using anhydrous magnesium sulfate, and the solvent was distilled out under reduced pressure. The residue was refined using alumina column chromatography (development solvent; ethyl acetate), and pulverized using isopropyl ether, to give the titled compound (13 mg) as a white powder.

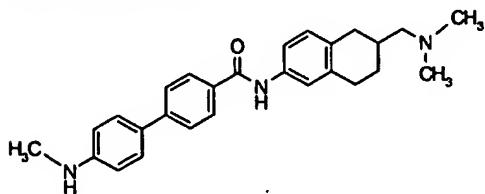
¹H-NMR (DMSO-d₆) δ : 1.32 (1H, m), 1.90 (2H, m), 2.15 (6H, s), 2.15-2.35 (3H, m), 2.77 (3H, m), 2.97 (6H, s), 6.82 (2H, d, J=8.4 Hz), 7.03 (1H, d, J=8.4 Hz), 7.48 (1H, d, J=8.1 Hz), 7.53 (1H, s), 7.63 (2H, d, J=8.7 Hz), 7.74 (2H, d, J=7.8 Hz), 7.98 (2H, d, J=8.4 Hz), 10.04 (1H, s).

FABMS(pos) 428.2[M+H]⁺

Melting point: 212 - 213°C (crystallization solvent: ethyl acetate-diisopropyl ether)

Example 38

N-[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-(methylamino)[1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as a white powder by the same method as in Example 37, using 4'-amino-N-[6-[(N,N-dimethyl)amino]methyl]-5,6,7,8-tetrahydro-2-naphthalenyl [1,1'-biphenyl]-4-carboxamide (150 mg, 0.375 mmol) obtained in Example 33, paraformaldehyde (15.0 mg, 0.50 mmol), and sodium cyanohydroborate (31.5 mg, 0.50 mmol).

¹H-NMR (DMSO-d₆) δ : 1.32 (1H, m), 1.89 (2H, m), 2.15 (6H, s), 2.15-2.31 (3H, m), 2.72 (7H, m), 5.94 (1H, m), 6.64 (2H,

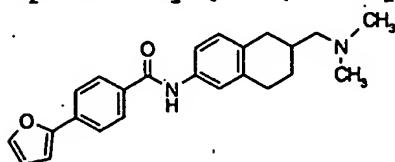
d, J=9.0 Hz), 7.03 (1H, d, J=8.7 Hz), 7.49 (4H, m), 7.70 (1H, d, J=8.4 Hz), 7.97 (2H, d, J=8.4 Hz), 10.02 (1H, s).
 FABMS(pos) 414.3[M+H]⁺

Melting point: 163 - 165°C (crystallization solvent: ethyl acetate-diisopropyl ether)

5

Example 39

N-[6-[(N,N-Dimethyl)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4-(2-furyl)benzamide



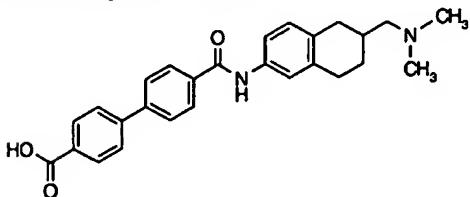
10

The titled compound (67 mg) was obtained as a white powder by the same method as in Example 16, using 4-bromo-N-[6-[(N,N-dimethyl)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide (250 mg, 0.645 mmol) obtained in Example 15, and 2-furylboric acid (86.7 mg, 0.775 mmol).
¹H-NMR (DMSO-d₆) δ: 1.40 (1H, m), 1.94 (2H, m), 2.25 (6H, s), 2.25-2.45 (3H, m), 2.88 (3H, m), 7.08 (1H, d, J=8.1 Hz), 7.26 (4H, m), 7.41 (1H, m), 7.60-7.74 (5H, m).
 FABMS(pos) 375.2[M+H]⁺

15

Example 40

4'-[[[6-[(N,N-Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]amino]carbonyl][1,1'-biphenyl]-4-carboxylic acid



20

Ethyl-4'-[[[6-[(N,N-dimethyl)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]amino]carbonyl][1,1'-biphenyl]-4-carboxylate (100 mg, 0.219 mmol) obtained in Example 35 was dissolved in a mixed solution of ethanol (3

25

ml) and water (0.5 ml). 1N aqueous sodium hydroxide solution (0.329 ml) was added to the reaction mixture at room temperature, which was stirred at 90°C for 5 hours.

After the solvent was distilled out under reduced pressure, water was added to the residue, then 1N hydrochloric acid (0.329 ml) was added and the reaction mixture was stirred. The precipitated crude product collected by filtration, and washed with water to give the titled compound (89 mg) as a white powder.

10 $^1\text{H-NMR}$ (DMSO-d₆) δ : 1.34 (1H, m), 1.91 (2H, m), 2.24 (6H, s), 2.24-2.30 (3H, m), 2.81 (3H, m), 7.05 (1H, d, J=8.4 Hz), 7.49 (1H, d, J=8.4 Hz), 7.55 (1H, s), 7.89 (4H, m), 8.07 (4H, m), 10.18 (1H, s).

Elemental analysis for C₂₇H₂₈N₂O₃ · 2H₂O

15 Calcd.: C, 69.81; H, 6.94; N, 6.03.

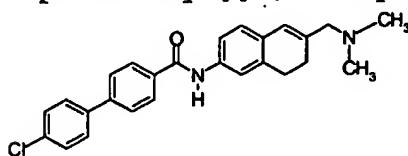
Found: C, 69.57; H, 7.01; N, 5.93.

Melting point: 143°C (decomposition) (crystallization solvent: water)

20

Example 41

4'-Chloro-N-[6-[(N,N-dimethyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



25 1) 6-Acetamido-1-tetralone (5.0 g, 0.0246 mol) synthesized according to a known method by documents (Journal of Organic Chemistry 27, 70 (1962)), was dissolved in 50 ml of DMF dimethylacetal, which was stirred at 110°C for 2 hours. The precipitate was collected by filtration, 30 and washed with ethyl acetate to give 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone (4.98 g) as a yellow powder.

$^1\text{H-NMR}$ (CDCl₃) δ : 2.19 (3H, s), 2.79-2.83 (2H, m), 2.88-

2.92 (2H, m), 3.11 (6H, s), 7.14-7.17 (1H, m), 7.68 (1H, s), 7.69 (1H, s), 7.95 (1H, d, J=8.1Hz), 7.96 (1H, s).

Melting point: 207 - 210°C (crystallization solvent: ethyl acetate)

5 2) The obtained 6-acetamido-2-(N,N-dimethylaminomethylidene)-1-tetralone (4.50 g, 0.0173 mol) was dissolved in methanol (50 ml), and sodium borohydride (6.56 g, 0.173 mol) was added to the solution under ice-cooling, which was stirred for 2 hours. The
10 reaction mixture was concentrated. Ethyl acetate and sodium hydrogencarbonate solution were added to the residue, and extraction was conducted. The ethyl acetate layer was concentrated, and 30 ml of tetrahydrofuran and 30 ml of 2N hydrochloric acid were added to the residue, which was refluxed with heating for 16 hours. The reaction mixture was concentrated, and ethyl acetate and 2N sodium hydroxide solution were added, and extraction was conducted. The ethyl acetate layer was concentrated, and the residue was refined using alumina column chromatography
15 (development solvent; ethyl acetate:n-hexane = 30:70), to give 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthaleneamine (1.60 g) as a colorless oily substance.
20 $^1\text{H-NMR}$ (CDCl_3) δ : 2.23 (6H, s), 2.28 (2H, t, J=8.4Hz), 2.74 (2H, t, J=8.4Hz), 2.95 (2H, s), 3.57-3.72 (2H, m), 6.25 (1H, s), 6.46-6.48 (2H, m), 6.83 (1H, d, J=8.7Hz).
25

3) The titled compound (1.12 g) was obtained as a white powder by the same method as in Example 1, using the obtained 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthaleneamine (1.00 g, 0.005 mol), and 4-chlorobiphenyl
30 carboxylic acid (2.31 g, 0.01 mol).

35 $^1\text{H-NMR}$ (CDCl_3) δ : 2.25 (6H, s), 2.34 (2H, t, J=7.8Hz), 2.86 (2H, t, J=7.8Hz), 2.99 (2H, s), 6.34 (1H, s), 7.03 (1H, d, J=8.7Hz), 7.39 (1H, d, J=8.1 Hz), 7.45 (2H, d, J=8.7), 7.48 (1H, s), 7.56 (2H, d, J=8.4 Hz), 7.67 (2H, d, J=8.4 Hz), 7.78 (1H, s), 7.94 (2H, d, J=8.4 Hz).

Elemental analysis for $\text{C}_{26}\text{H}_{25}\text{ClN}_2\text{O}$

Calcd. : C, 74.90; H, 6.04; N, 6.72.

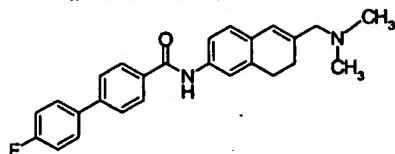
Found : C, 74.64; H, 6.14; N, 6.56.

Melting point: 204 - 207°C (crystallization solvent: ethyl acetate - n-hexane)

5

Example 42

4'-Fluoro-N-[6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



10 The titled compound (990 mg) was obtained as a white powder by the same method as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine (936 mg, 4.62 mmol) obtained in Example 41-2), and 4-fluorobiphenylcarboxylic acid (1.00 g, 4.62 mmol).

15 $^1\text{H-NMR}$ (CDCl_3) δ : 2.25 (6H, s), 2.34 (2H, t, $J=8.1\text{Hz}$), 2.85 (2H, t, $J=8.1\text{Hz}$), 2.99 (2H, s), 6.34 (1H, s), 7.02 (1H, d, $J=8.1\text{Hz}$), 7.13-7.19 (2H, m), 7.38-7.41 (1H, m), 7.48 (1H, s), 7.56-7.61 (2H, m), 7.65 (2H, d, $J=8.4\text{ Hz}$), 7.80 (1H, s), 7.93 (2H, d, $J=8.5\text{Hz}$).

20 Elemental analysis for $\text{C}_{26}\text{H}_{25}\text{FN}_2\text{O}$

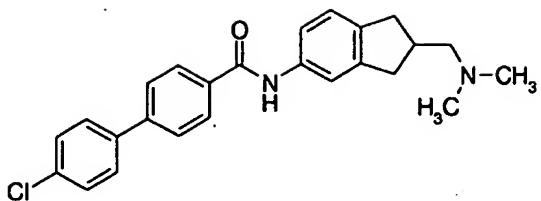
Calcd. : C, 77.97; H, 6.29; N, 6.99.

Found : C, 77.90; H, 6.23; N, 6.58.

25 Melting point: 190 - 193°C (crystallization solvent: ethyl acetate - n-hexane)

Example 43

4'-Chloro-N-[2-[(dimethylamino)methyl]-2,3-dihydro-1H-inden-5-yl][1,1'-biphenyl]-4-carboxamide



Concentrated hydrochloric acid (1 ml) was added to N-[2-[(dimethylamino)methyl]-2,3-dihydro-1H-inden-5-yl]acetamide (48.9 mg, 0.210 mmol) obtained in Reference Example 48, which was stirred at 110°C for 2 hours, and the solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with potassium carbonate solution and saturated aqueous sodium chloride solution, dried using anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. Using the oily substance obtained, the same operation as in Example 1 was conducted to give the titled compound (30 mg).

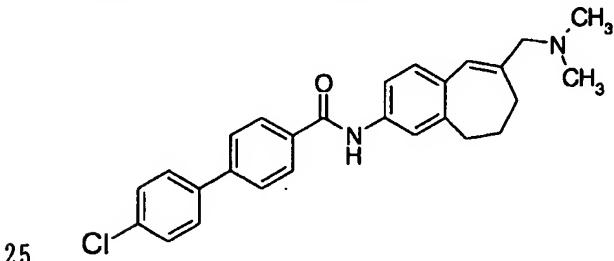
¹H NMR (DMSO-d₆) δ: 2.16 (6H, s), 2.22 (2H, d, J = 6.7 Hz), 2.61 (4H, m), 2.97 (1H, m), 7.15 (1H, d, J = 8.1 Hz), 7.47 (1H, d, J = 8.1 Hz), 7.56 (2H, d, J = 8.4 Hz), 8.05 (2H, d, J = 8.4 Hz), 10.17 (1H, s).

FAB(pos) 405.1 [M+H]⁺

Melting point: 192 - 194°C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 44

4'-Chloro-N-[8-[(dimethylamino)methyl]-6,7-dihydro-5H-benzo[a]cyclohepten-3-yl][1,1'-biphenyl]-4-carboxamide



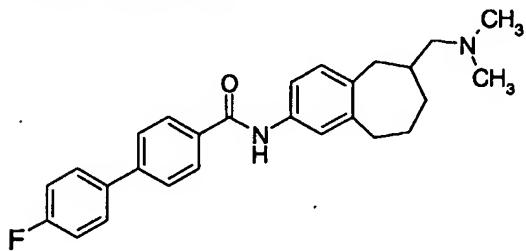
The titled compound was obtained by carrying out the same operation as in Example 1, using 8-

[(dimethylamino)methyl]-6,7-dihydro-5H-benzo[a]cyclohepten-3-amine obtained in Reference Example 50.

¹H-NMR (CDCl₃) δ : 1.96-2.10 (2H, m), 2.25 (6H, s), 2.39 (2H, t, J = 6.4 Hz), 2.79-2.85 (2H, m), 2.96 (2H, s), 6.40 (1H, s), 7.15 (1H, d, J = 8.6Hz), 7.40-7.52 (4H, m), 7.56 (2H, d, J = 8.4Hz), 7.67 (2H, d, J = 8.1Hz), 7.81 (1H, s), 7.94 (2H, d, J = 8.1 Hz).

Melting point: 183-185°C (crystallization solvent: ethyl acetate - diethyl ether)

Example 45
4'-Fluoro-N-[6-[(dimethylamino)methyl]-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-2-yl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(dimethylamino)methyl]-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-2-amine obtained in Reference Example 51.

¹H-NMR (CDCl₃) δ : 1.40-1.68 (3H, m), 1.85-2.20 (10H, m), 2.55-2.92 (4H, m), 7.13-7.20 (3H ,m), 7.35-7.43 (2H, m), 7.56-7.67 (4H, m), 7.77 (1H, s), 7.93 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₇H₂₉FN₂O

Calcd. : C, 77.85; H, 7.02; N, 6.73.

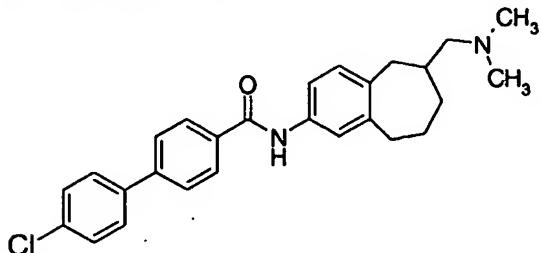
Found : C, 78.18; H, 7.09; N, 6.74.

Melting point: 167 - 169°C (crystallization solvent: diethyl ether)

Example 46

4'-Chloro-N-[6-[(dimethylamino)methyl]-6,7,8,9-

tetrahydro-5H-benzo[a]cyclohepten-2-yl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the
5 same operation as in Experiment Example 1, using 6-[
[(dimethylamino)methyl]-6,7,8,9-tetrahydro-5H-
benzo[a]cyclohepten-2-amine obtained in Reference Example
51.

¹H-NMR (CDCl₃) δ : 1.40-1.67 (3H, m), 1.85-2.20 (10H, m),
10 2.55-2.92 (4H, m), 7.15 (1H, d, J = 8.1 Hz), 7.35-7.46 (4H,
m), 7.56 (2H, d, J = 8.4 Hz), 7.66 (2H, d, J = 8.1 Hz), 7.77
(1H, s), 7.93 (2H, d, J = 8.4 Hz).

Elemental analysis for C₂₇H₂₉ClN₂O

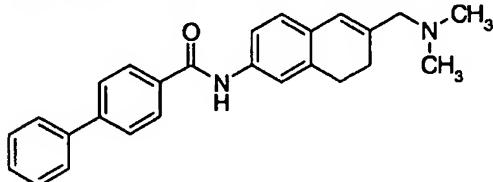
Calcd. : C, 74.90; H, 6.75; N, 6.47.

15 Found: C, 74.77; H, 6.65; N, 6.43.

Melting point: 173 - 175°C (crystallization solvent:
diethyl ether)

Example 47

20 N-[6-[(Dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the
same operation as in Example 1, using 6-[(N,N-
25 dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine
obtained in Example 41-2).

¹H NMR (CDCl₃) δ : 2.25 (6H, s), 2.33 (2H, t, J = 5.4 Hz),

2.84 (2H, t, J = 5.4 Hz), 2.98 (2H, s), 6.34 (1H, s), 7.01 (1H, d, J = 7.8 Hz), 7.32-7.94 (12H, m).

Elemental analysis for C₂₆H₂₆N₂O

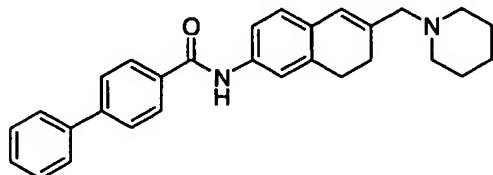
Calcd. : C, 81.64; H, 6.85; N, 7.32.

5 Found: C, 81.65; H, 6.79; N, 6.91.

Melting point: 173 - 175°C (crystallization solvent:
tetrahydrofuran - n-hexane)

Example 48

10 N-[6-(1-Piperidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



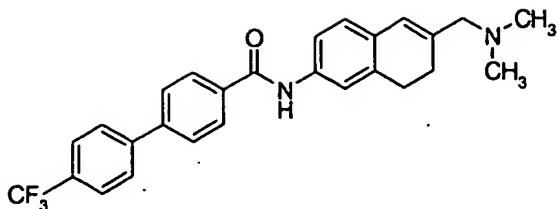
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 52.

¹H NMR (CDCl₃) δ : 1.46-1.59 (6H, m), 2.31-2.36 (6H, m), 2.84 (2H, t, J = 8.0 Hz), 3.02 (2H, s), 6.34 (1H, s), 7.02 (1H, d, J = 8.1 Hz), 7.37-7.50 (4H, m), 7.63 (2H, d, J = 6.9 Hz), 20 7.71 (2H, d, J = 8.1 Hz), 7.79 (1H, s), 7.94 (2H, d, J = 8.1 Hz).

Melting point: 156 - 158°C (crystallization solvent:
tetrahydrofuran - n-hexane)

25 Example 49

N-[6-[(Dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]-4'-trifluoromethyl[1,1'-biphenyl]-4-carboxamide



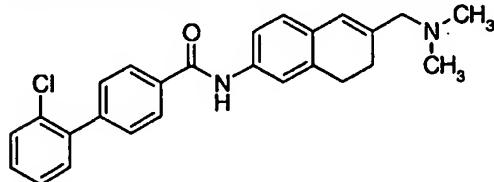
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Example 41-2).

5 ¹H NMR (CDCl₃) δ : 2.25 (6H, s), 2.34 (d, J = 5.1 Hz), 2.86 (2H, d, J = 5.1 Hz), 2.99 (2H, s), 6.35 (1H, s), 7.04 (1H, d, J = 8.4 Hz), 7.40 (1H, d, J = 3.3 Hz), 7.49 (1H, s), 7.70-7.79 (6H, m), 7.87 (2H, d, J = 8.4 Hz).

10 Melting point: 214 - 216°C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 50

15 2'-Chloro-N-[6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Example 41-2).

20 ¹H NMR (CDCl₃) δ : 2.25 (6H, s), 2.34 (d, J = 5.1 Hz), 2.85 (2H, d, J = 5.1 Hz), 3.00 (2H, s), 6.34 (1H, s), 6.69 (1H, s), 7.02 (1H, d, J = 8.4 Hz), 7.31-7.57 (8H, m), 7.85 (1H, s), 7.92 (2H, d, J = 7.8 Hz).

25 Elemental analysis for C₂₆H₂₅ClN₂O

Calcd. : C, 74.90; H, 6.04; N, 6.72

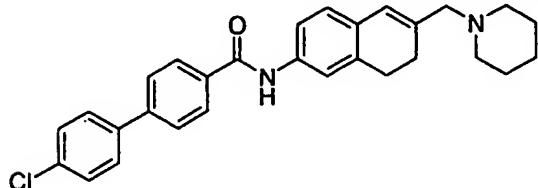
Found : C, 74.49; H, 5.65; N, 6.06.

Melting point: 145 - 147°C (crystallization solvent: ethyl

acetate - n-hexane)

Example 51

4'-Chloro-N-[6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



After N,N-dimethylformaldehyde solution (5 ml) of 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide (225 mg) obtained in Reference Example 56, piperidine (0.16 ml), and diisopropylethylamine (0.282 ml) was stirred at room temperature for 15 hours, which was heated at 120°C for 2 hours. The residue obtained by concentrating the reaction mixture was dissolved in water-ethyl acetate, then extracted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, dried using anhydrous magnesium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was refined using alumina column chromatography (development solvent; tetrahydrofuran:n-hexane = 1:5), and crystallized using tetrahydrofuran - n-hexane to give the titled compound (110 mg).

¹H NMR (CDCl₃) δ: 1.26-1.61 (6H, m), 2.30-2.36 (6H, m), 2.83 (2H, t, J = 8.4 Hz), 3.02 (2H, s), 6.33 (1H, s), 7.01 (1H, d, J = 8.1 Hz), 7.36-7.49 (4H, m), 7.55 (2H, d, J = 8.4 Hz), 7.66 (2H, d, J = 8.4 Hz), 7.81 (1H, s), 7.93 (2H, d, J = 8.1 Hz).

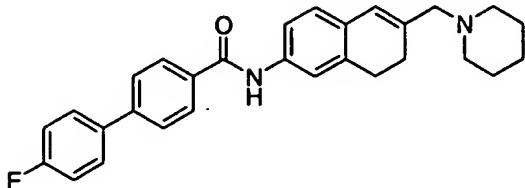
Melting point: 209 - 211°C (crystallization solvent:
tetrahydrofuran - n-hexane)

30

Example 52

4'-Fluoro-N-[6-(1-piperidinylmethyl)-7,8-dihydro-2-

naphthalenyl][1,1'-biphenyl]4-carboxamide



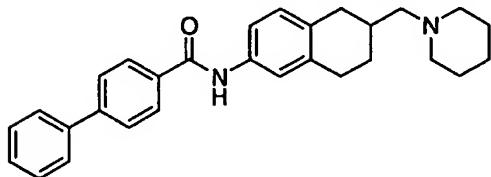
5 The titled compound was obtained by carrying out the same operation as in Example 1, using the 6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalene amine obtained in Reference example 52.

10 ¹H NMR (CDCl₃) δ : 1.45-1.58 (6H, m), 2.29-2.37 (6H, m), 2.82 (2H, t, J = 8.0 Hz), 3.01 (2H, s), 6.33 (1H, s), 6.98-7.93 (12H, m).

15 Melting point: 190 - 192°C (crystallization solvent: tetrahydrofuran - n-hexane)

Example 53

15 N-[6-(1-Piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



20 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

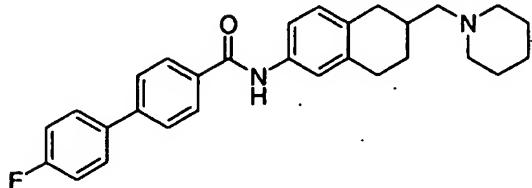
25 ¹H NMR (CDCl₃) δ : 1.37-1.60 (8H, m), 1.96-2.00 (2H, m), 2.24-2.44 (5H, m), 2.82-2.93 (3H, m), 7.09 (1H, d, J = 8.3 Hz), 7.30-7.33 (1H, m), 7.38-7.65 (6H, m), 7.70 (2H, d, J = 8.4 Hz), 7.76 (1H, s), 7.93 (2H, d, J = 8.4 Hz).

25 Melting point: 160 - 162°C (crystallization solvent: tetrahydrofuran - n-hexane)

Example 54

4'-Fluoro-N-[6-(1-piperidinylmethyl)-5,6,7,8-

tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

¹H NMR (CDCl₃) δ: 1.36-1.52 (8H, m), 2.29-2.31 (2H, m), 2.24-2.45 (6H, m), 2.82-2.93 (3H, m), 7.08-7.33 (4H, m), 7.44 (1H, s), 7.57-7.66 (4H, m), 7.74 (1H, s), 7.92 (2H, 10 J = 8.1 Hz).

Elemental analysis for C₂₉H₃₁FN₂O

Calcd.: C, 78.70; H, 7.08; N, 6.33.

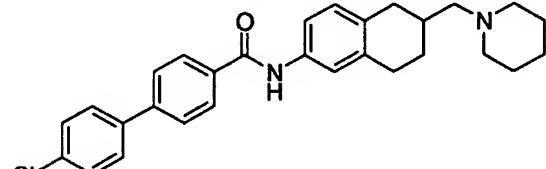
Found: C, 78.40; H, 7.09; N, 6.09.

Melting point: 179 - 181°C (crystallization solvent: ethyl acetate)

Example 55

4'-Chloro-N-[6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

20



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

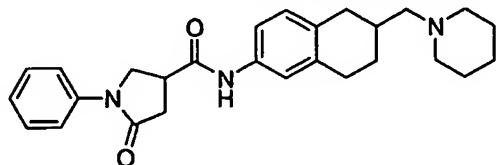
25

¹H NMR (CDCl₃) δ: 1.25-1.71 (8H, m), 1.95-2.00 (2H, m), 2.25-2.45 (6H, m), 2.83-2.93 (3H, m), 7.09 (1H, d, J = 8.3 Hz), 7.30-7.32 (1H, m), 7.43-7.45 (3H, m), 7.55 (2H, d, J = 8.1 Hz), 7.65 (2H, d, J = 8.4 Hz), 7.77 (1H, s), 7.93 (2H, d, J = 8.1 Hz).

Melting point: 202 - 203°C (crystallization solvent:
tetrahydrofuran - n-hexane)

Example 56

5 5-Oxo-1-phenyl-N-[6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl]-3-pyrrolidinecarboxamide



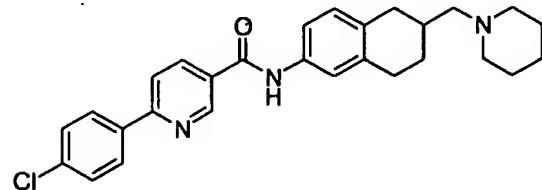
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

¹H NMR (CDCl₃) δ: 1.03-3.33(22H, m), 3.97 (1H, t, J = 8.4 Hz), 4.21 (1H, dd, J = 6.8, 7.1 Hz), 6.91-7.63 (9H, m). Elemental analysis for C₂₇H₃₃N₃O₂
15 Calcd.: C, 75.14; H, 7.71; N, 9.74.
Found: C, 75.01; H, 7.33; N, 9.43.

Melting point: 162 - 164°C (crystallization solvent: ethyl acetate)

20 **Example 57**

6-(4-Chlorophenyl)-N-[6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl]nicotinamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

¹H NMR (CDCl₃) δ: 1.30-2.40 (16H, m), 2.82-2.92 (3H, m), 7.09 (1H, d, J = 8.1 Hz), 7.26-7.48 (4H, m), 7.80 (2H, d,

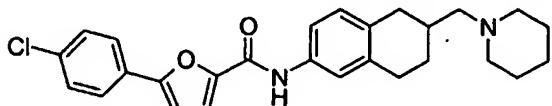
J = 8.7 Hz), 7.99 (2H, d, *J* = 8.7 Hz), 8.23 (d, 1H, *J* = 6.3 Hz), 9.11 (1H, s).

Melting point: 193 - 195°C (crystallization solvent: ethyl acetate)

5

Example 58

5-(4-Chlorophenyl)-N-[6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl]-2-furamide

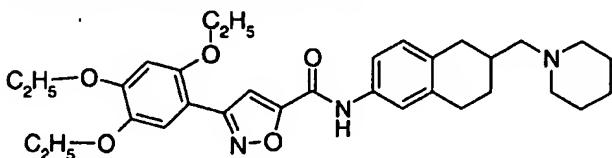


10 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

15 ^1H NMR (CDCl_3) δ : 1.23-1.61 (7H, m), 1.96-2.00 (2H, m), 2.24-2.43 (7H, m), 2.80-2.92 (3H, m), 6.75 (1H, d, *J* = 3.6 Hz), 7.07 (1H, d, *J* = 8.4 Hz), 7.27 (1H, d, *J* = 3.6 Hz), 7.32-7.42 (4H, m), 7.66 (2H, d, *J* = 8.4 Hz), 8.32 (1H, s).

Example 59

20 N-[6-(1-Piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl]-3-(2,4,5-triethoxyphenyl)-5-isoxazolecarboxamide

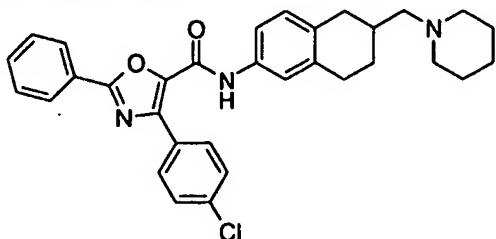


25 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

30 ^1H NMR (CDCl_3) δ : 1.42-1.60 (18H, m), 1.97-2.36 (7H, m), 2.80-2.95 (3H, m), 4.06-4.18 (6H, m), 6.58 (1H, s), 7.09 (1H, d, *J* = 8.4 Hz), 7.35 (1H, d, *J* = 8.1 Hz), 7.44 (1H, s), 7.50 (1H, s), 7.55 (1H, s), 8.16 (1H, s).

Example 60

4-(4-Chlorophenyl)-2-phenyl-N-[6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl]-1,3-oxazole-5-carboxamide

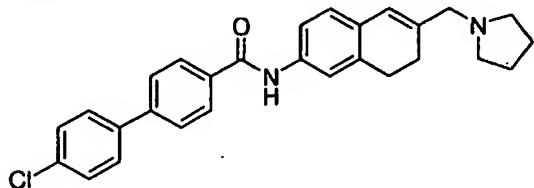


The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-piperidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 53.

¹H NMR (CDCl₃) δ: 1.26-1.58 (7H, m), 1.90-2.00 (2H, m), 2.22-2.35 (7H, m), 2.70-2.95 (3H, m), 7.06 (1H, d, J = 8.1 Hz), 7.25-7.51 (7H, m), 8.04-8.32 (5H, m).

15 Example 61

4'-Chloro-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.

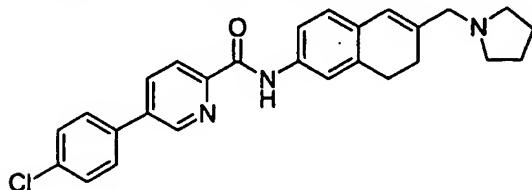
Melting point: 185 - 187°C (crystallization solvent:
tetrahydrofuran - n-hexane)

¹H NMR (CDCl₃) δ: 1.83 (4H, s), 2.35 (2H, t, J = 8.1 Hz), 2.52 (4H, s), 2.84 (2H, t, J = 8.1 Hz), 3.18 (2H, s), 6.36 (1H, s), 7.02 (1H, d, J = 8.4 Hz), 7.39-7.56 (6H, m), 7.66

(2H, d, J = 7.5 Hz), 7.82 (1H, s), 7.93 (2H, d, J = 7.5 Hz).

Example 62

5-(4-Chlorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-2-pyridinecarboxamide

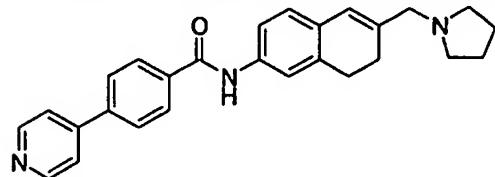


The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.

¹H NMR (CDCl₃) δ: 1.80 (6H, s), 2.37 (2H, t, J = 8.1 Hz), 2.52 (4H, s), 2.87 (2H, t, J = 8.1 Hz), 3.18 (2H, s), 6.37 (1H, s), 7.03 (1H, d, J = 7.8 Hz), 7.48-7.61 (6H, m), 8.04 (1H, dd, J = 8.1, 2.1 Hz), 8.35 (1H, d, J = 8.1 Hz), 8.78 (1H, s), 9.95 (1H, s).

Example 63

4-(4-Pyridinyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]benzamide

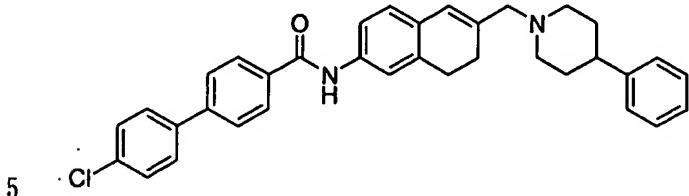


The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H NMR (CDCl₃) δ: 1.79-1.83 (6H, m), 2.35 (2H, t, J = 8.1 Hz), 2.53 (4H, s), 2.73 (2H, t, J = 8.1 Hz), 3.18 (2H, s), 6.36 (1H, s), 7.02 (1H, d, J = 7.8 Hz), 7.38 (1H, d, J = 8.1 Hz), 7.48 (1H, s), 7.71-7.78 (4H, m), 7.89 (1H, s), 7.99 (1H, d, J = 8.4 Hz), 8.32 (2H, d, J = 8.4 Hz).

Example 64

4'-Chloro-N-[6-[(4-phenyl-1-piperidinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



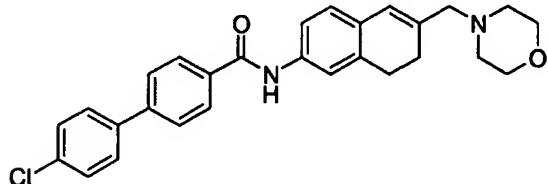
The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.

10 ¹H NMR (CDCl₃) δ: 1.83-2.10 (6H, m), 2.37 (2H, t, J = 8.1 Hz), 2.47-2.54 (1H, m), 2.86 (2H, t, J = 8.1 Hz), 3.03-3.10 (2H, m), 3.10 (2H, s), 6.37 (1H, s), 7.03 (1H, d, J = 8.4 Hz), 7.19-7.57 (11H, m), 7.66 (2H, d, J = 8.4 Hz), 7.81 (1H, s), 7.94 (2H, d, J = 8.4 Hz).

15 Melting point: 228 - 230°C (crystallization solvent:
tetrahydrofuran - n-hexane)

Example 65

4'-Chloro-N-[6-(4-morpholinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



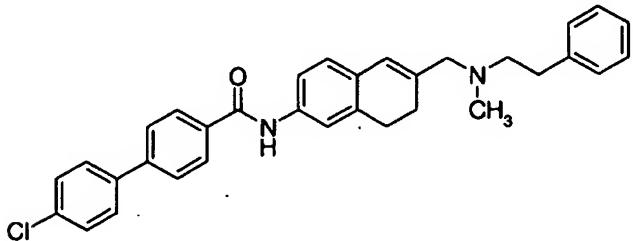
The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.

25 ¹H NMR (CDCl₃) δ: 2.34 (2H, t, J = 7.8 Hz), 2.45 (4H, s), 2.84 (2H, t, J = 7.8 Hz), 3.06 (2H, s), 3.73 (4H, s), 6.36 (1H, s), 7.02 (1H, d, J = 8.1 Hz), 7.36-7.57 (6H, m), 7.67 (2H, d, J = 8.4 Hz), 7.80 (1H, s), 7.94 (2H, d, J = 8.4 Hz).

Melting point: 194 - 195°C (crystallization solvent:
tetrahydrofuran - n-hexane)

Example 66

5 4'-Chloro-N-(6-[[methyl(2-phenylethyl)amino]methyl]-
7,8-dihydro-2-naphthalenyl[1,1'-biphenyl]-4-carboxamide

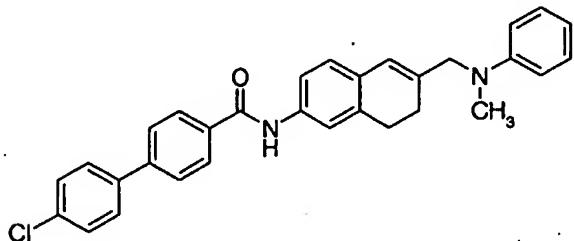


The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.
10 ^1H NMR (CDCl_3) δ : 2.25-2.32 (2H, m), 2.32 (3H, s), 2.60-2.66 (2H, m), 2.77-2.83 (4H, m), 3.10 (2H, s), 6.32 (1H, s), 6.93-7.95 (16H, m).

15 Melting point: 173 - 175°C (crystallization solvent:
tetrahydrofuran - n-hexane)

Example 67

4'-Chloro-N-[6-[methylanilino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide
20



The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.
25 ^1H NMR (CDCl_3) δ : 2.20-2.30 (2H, m), 2.25 (3H, s), 2.85-2.90

(2H, m), 3.00 (2H, s), 6.30 (1H, s), 6.74-7.95 (146H, m).
 Melting point: 177 - 179°C (crystallization solvent:
 tetrahydrofuran - n-hexane)

5 Example 68

4'-Chloro-N-[6-[(4-phenyl-1-piperadinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

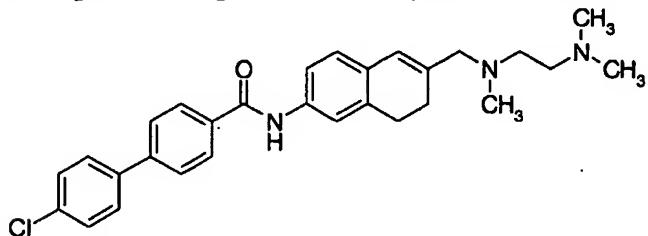


10 The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.
¹H NMR (CDCl₃) δ: 2.37 (2H, t, J = 8.1 Hz), 2.62 (4H, s), 2.86 (2H, t, J = 8.4 Hz), 3.13 (2H, s), 3.22 (4H, s), 6.39 (1H, s), 6.85-7.95 (16H, m).

Melting point: 228 - 230°C (crystallization solvent:
 tetrahydrofuran - n-hexane)

Example 69

20 4'-Chloro-N-[6-[[[2-(dimethylamino)ethyl](methyl)amino]methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



25 The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.
¹H NMR (CDCl₃) δ: 2.25 (6H, s), 2.26 (3H, s), 2.33 (2H, t,

$J = 8.1$ Hz), 2.44-2.50 (4H, m), 2.84 (2H, t, $J = 8.1$ Hz), 3.07 (2H, s), 6.35 (1H, s), 7.02 (1H, d, $J = 8.4$ Hz), 7.37-7.57 (6H, m), 7.67 (2H, d, $J = 8.1$ Hz), 7.80 (1H, s), 7.94 (2H, d, $J = 8.4$ Hz).

5 Melting point: 156 - 158°C (crystallization solvent: tetrahydrofuran - n-hexane)

Example 70

10 4'-Fluoro-N-[6-(4-morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



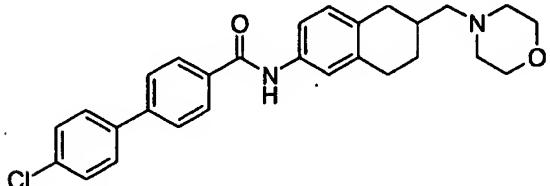
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(4-morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 57.

15 ^1H NMR (CDCl_3) δ : 1.40-1.50 (1H, m), 1.90-2.10 (2H, m), 2.29-2.45 (7H, m), 2.80-2.92 (3H, m), 3.72-3.75 (4H, m), 7.07-7.33 (4H, m), 7.46 (1H, s), 7.56-7.66 (4H, m), 7.78 (1H, s), 7.92 (2H, d, $J = 8.1$ Hz).

20 Melting point: 188 - 190°C (crystallization solvent: ethyl acetate)

Example 71

25 4'-Chloro-N-[6-(4-morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(4-morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine

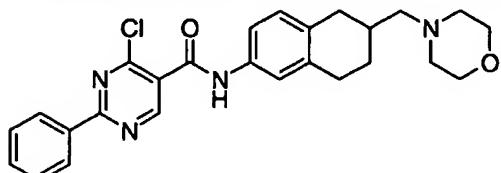
obtained in Reference Example 57.

^1H NMR (CDCl_3) δ : 1.40-1.50 (1H, m), 1.90-2.10 (2H, m), 2.32-2.45 (7H, m), 2.80-2.90 (3H, m), 3.70-3.80 (4H, m), 7.10-7.92 (12H, m).

5 Melting point: 216 - 218°C (crystallization solvent: ethyl acetate)

Example 72

10 4-Chloro-N-[6-(4-morpholinylmethyl)-5,6,7,8-tetrahydro-
2-naphthalenyl]-2-phenyl-5-pyrimidinecarboxamide

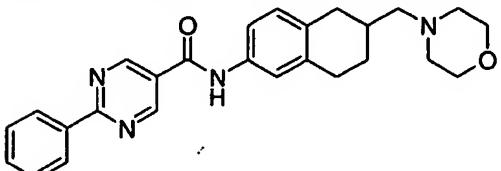


The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(4-morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 57.

15 ^1H NMR (CDCl_3) δ : 1.40-1.50 (1H, m), 1.95-2.05 (2H, m), 2.29-2.45 (7H, m), 2.80-2.95 (3H, m), 3.73 (4H, t, J = 4.5 Hz), 7.10 (1H, d, J = 8.1 Hz), 7.32 (1H, d, J = 8.1 Hz), 7.42 (1H, s), 7.49-7.56 (3H, m), 8.25 (1H, s), 8.48 (2H, d, J = 6.6 Hz), 9.20 (1H, s)

Example 73

20 N-[6-(4-Morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl]-2-phenyl-5-pyrimidinecarboxamide



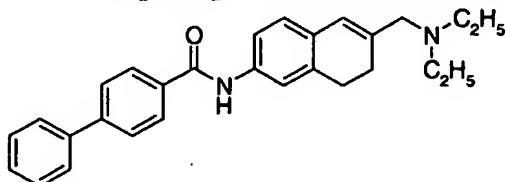
25 The titled compound was obtained by carrying out the same operation as in Reference Example 48, using 4-chloro-N-[6-(4-morpholinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl]-2-phenyl-5-pyrimidinecarboxamide obtained in

Example 72.

¹H NMR (CDCl₃) δ : 1.21-1.30 (1H, m), 1.93-2.03 (2H, m), 2.28
 -2.44 (7H, m), 2.80-2.90 (3H, m), 3.73 (4H, t, J = 4.8 Hz),
 7.07 (1H, d, J = 8.1 Hz), 7.26 -7.30 (1H, m), 7.39 (1H, s),
 5 7.51-7.53 (3H, m), 8.00 (1H, s), 8.50 (2H, dd, J = 8.1, 2.4
 Hz), 9.21 (2H, s)

Example 74

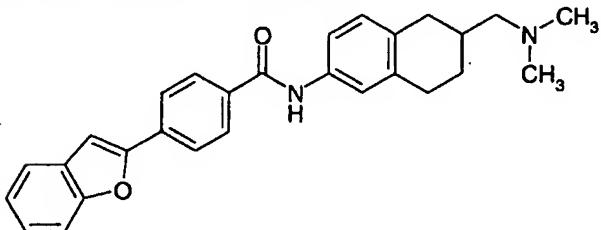
N-[6-[(Diethylamino)methyl]-7,8-dihydro-2-naphthalenyl]
 10 [1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 51, using N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 58.
 15 ¹H NMR (CDCl₃) δ : 1.24 (6H, t, J = 7.2 Hz), 2.33 (2H, t, J = 5.1 Hz), 2.53 (4H, q, J = 7.2 Hz), 2.84 (2H, t, J = 5.1 Hz), 3.11 (2H, s), 6.36 (1H, s), 7.02 (1H, d, J = 8.1 Hz), 7.37-7.50 (5H, m), 7.63 (2H, d, J = 8.7 Hz), 7.71 (2H, d, J = 8.4 Hz), 20 7.79 (1H, s), 7.93 (2H, d, J = 8.4 Hz).
 Melting point: 153 - 155°C (crystallization solvent:
 tetrahydrofuran - n-hexane)

Example 75

25 4-(2-Benzo[b]furanyl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide



The titled compound was obtained by carrying out the

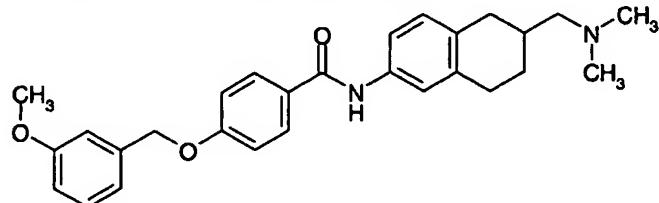
same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

Melting point: 192 - 194°C (crystallization solvent: tetrahydrofuran-isopropyl ether)

5

Example 76

4-(3-Methoxybenzyloxy)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide



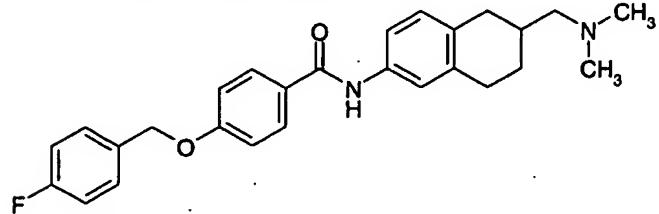
10 The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

Melting point: 102 - 104°C (crystallization solvent: isopropyl ether)

15

Example 77

4-(4-Fluorobenzyloxy)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide



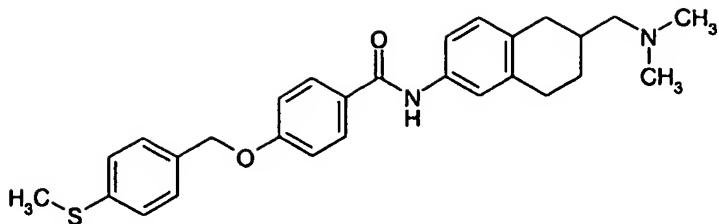
20 The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

Melting point: 165 - 167°C (crystallization solvent: tetrahydrofuran-hexane)

25

Example 78

4-[4-(Methylsulfanyl)benzyloxy]-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide

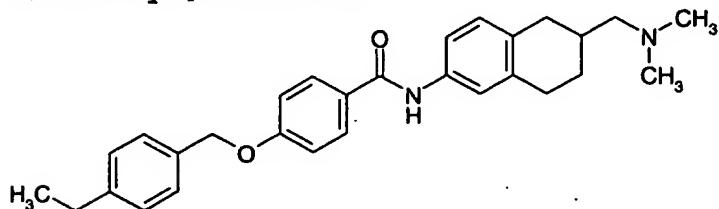


The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

5 Melting point: 162 - 163°C (crystallization solvent:
tetrahydrofuran-hexane)

Example 79

10 4-(4-Ethylbenzyloxy)-N-[2-(N,N-dimethylamino)methyl]-6-tetralinylbenzamide

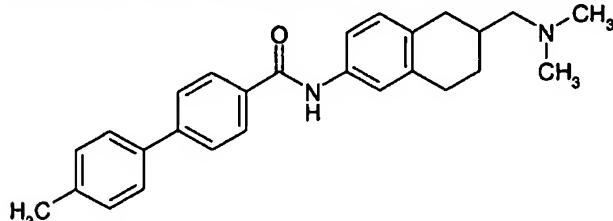


The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

15 Melting point: 120 - 122°C (crystallization solvent:
tetrahydrofuran-isopropyl ether)

Example 80

20 (4'-Methylbiphenyl-4-yl)-N-[2-(N,N-dimethylamino)methyl]-6-tetralinyl]carboxamide



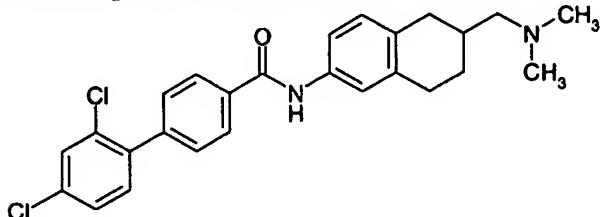
The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-

dimethylamino)methyl]tetralin hydrochloride.

Melting point: 181 - 182°C (crystallization solvent: ethyl acetate-hexane)

5 Example 81

(2',4'-Dichlorobiphenyl-4-yl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]carboxamide

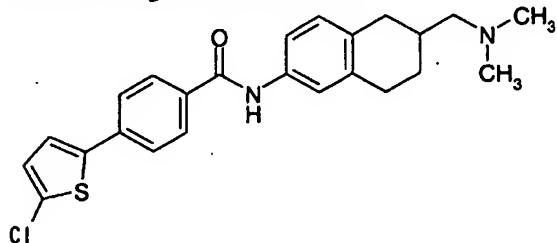


The titled compound was obtained by carrying out the
10 same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

Melting point: 188 - 189°C (crystallization solvent:
tetrahydrofuran-hexane)

15 Example 82

4-(5-Chloro-2-thienyl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]benzamide

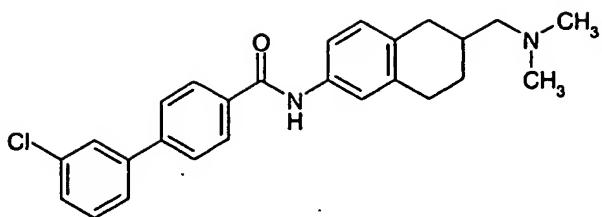


The titled compound was obtained by carrying out the
20 same operation as in Example 4, using 6-amino-2-(N,N-dimethylamino)methyltetraline hydrochloride.

Melting point: 167 - 169°C (crystallization solvent: ethyl acetate-hexane)

25 Example 83

(3'-Chlorobiphenyl-4-yl)-N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]carboxamide

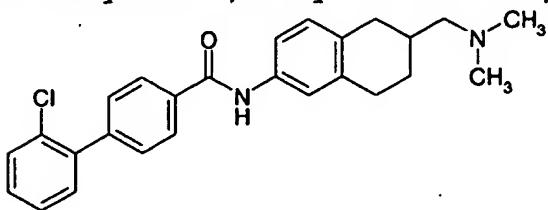


The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

5 Melting point: 138 - 139°C (crystallization solvent:
tetrahydrofuran-isopropyl ether)

Example 84

10 (2'-Chlorobiphenyl-4-yl)-N-[2-(N,N-dimethylamino)methyl]-6-tetralinyl]carboxamide

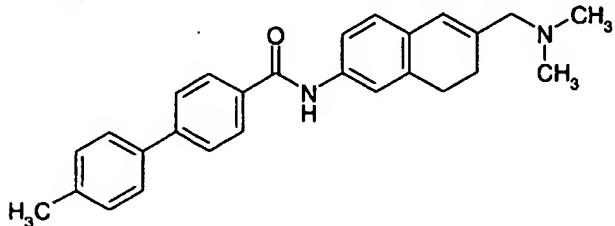


The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

15 Melting point: 176 - 177°C (crystallization solvent:
tetrahydrofuran-hexane)

Example 85

20 4'-Methyl-N-[6-(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl[1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine

obtained in Example 41-2).

¹H-NMR (CDCl₃) δ : 2.25 (6H, s), 2.33 (2H, t, J = 8.1 Hz), 2.41 (3H, s), 2.84 (2H, t, J = 8.1 Hz), 2.98 (2H, s), 6.33 (1H, s), 7.01 (1H, d, J = 7.8 Hz), 7.39 (1H, d, J = 8.4 Hz), 7.48 (1H, s), 7.52 (2H, d, J = 7.8 Hz), 7.67 (2H, d, J = 8.1 Hz), 7.84 (1H, s), 7.91 (2H, d, J = 8.1 Hz).

Elemental analysis for C₂₇H₂₈N₂O

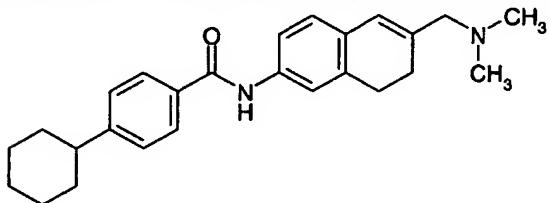
Calcd.: C, 81.78; H, 7.12; N, 7.06

Found: C, 81.51; H, 7.22; N, 6.93

10 Melting point: 195 - 196°C (crystallization solvent: ethyl acetate-diisopropyl ether)

Example 86

15 4-Cyclohexyl-N-[6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]benzamide



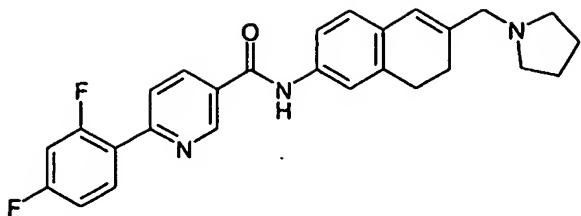
The titled compound was obtained by carrying out the same operation as in Example 1, using the 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Example 41-2.

20 ¹H-NMR (CDCl₃) δ : 1.20-1.52 (4H, m), 1.71-1.96 (6H, m), 2.25 (6H, s), 2.33 (2H, t, J = 8.1 Hz), 2.50-2.62 (1H, m), 2.84 (2H, t, J = 8.1 Hz), 2.99 (2H, s), 6.33 (1H, s), 7.00 (1H, d, J = 7.8 Hz), 7.31 (2H, d, J = 8.1 Hz), 7.36 (1H, d, J = 7.8 Hz), 7.46 (1H, brs), 7.75 (1H, s), 7.78 (2H, d, J = 8.1 Hz).

Melting point: 179 - 181°C (crystallization solvent: ethyl acetate-diisopropyl ether)

30 Example 87

6-(2,4-Difluorophenyl)-N-[6-[(1-pyrrolidinyl)methyl]-7,8-dihydro-2-naphthalenyl]nicotinamide



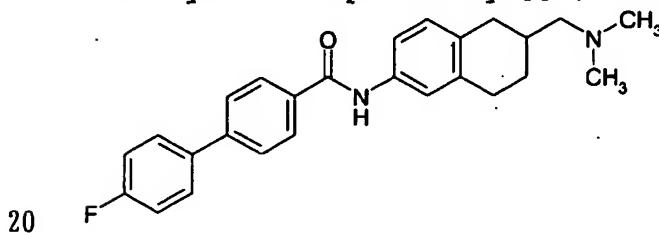
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ : 1.81 (4H, m), 2.37 (2H, t, J = 8.1 Hz), 2.54 (4H, m), 2.86 (2H, t, J = 8.1 Hz), 3.18 (2H, s), 6.37 (1H, s), 6.93 (1H, m), 7.04 (2H, m), 7.38 (1H, m), 7.47 (1H, s), 7.77 (1H, s), 7.91 (1H, m), 8.13 (1H, m), 8.24 (1H, m), 9.16 (1H, s).

Elemental analysis for C₂₇H₂₆F₂N₃O
Calcd.: C, 72.79; H, 5.66; N, 9.43
Found: C, 72.65; H, 5.52; N, 9.73
Melting point: 169 - 170°C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 88

4'-Fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 4, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

¹H-NMR (CDCl₃) δ : 1.41 (1H, m), 1.95 (2H, m), 2.25-2.45 (3H, m), 2.36 (6H, s), 2.85-2.94 (3H, m), 7.13 (3H, m), 7.30 (1H, m), 7.46 (1H, s), 7.59 (2H, m), 7.65 (2H, d, J = 8.1 Hz), 7.74 (1H, s), 7.93 (2H, d, J = 8.1 Hz).

Elemental analysis for C₂₆H₂₇FN₂O

Calcd.: C, 77.58; H, 6.76; N, 6.96

Found: C, 77.72; H, 6.49; N, 6.79

Melting point: 184 - 186°C (crystallization solvent: ethyl acetate - diisopropyl ether)

5

Example 89

(+) -4'-Fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide, and (-)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

Optical resolution of 4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide (2.00 g)

15 obtained in Example 88 was conducted by sample-splitting HPLC using a chiral column (Daicel Co., CHIRALCEL OD 500 mmD × 500 mmL; moving phase n-hexane:ethanol = 85:15), to give (+) form (1.00 g; 99.8%ee) and (-) form (0.89 g; >99.9%ee) as powders. The powders obtained were
20 respectively recrystallized using ethyl acetate - diisopropyl ether, to give the (+) form (855 mg) and (-) form (754 mg) of the titled compounds. The optical rotation of both compounds are shown below.

(+)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

Optical rotation: $[\alpha]_D = +50.8^\circ$ C=0.494% (methanol)

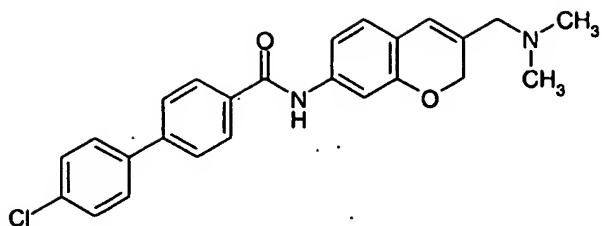
(-)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

Optical rotation: $[\alpha]_D = +51.2^\circ$ C=0 .492% (methanol)

30

Example 90

4'-Chloro-N-[3-[(N,N-dimethylamino)methyl]-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide



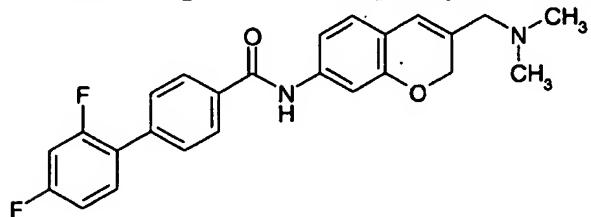
The titled compound was obtained by carrying out the same operation as in Example 1, using 3-[N,N-dimethylamino)methyl]-2H-chromen-7-amine obtained in Reference Example 59.

¹H-NMR (CDCl₃) δ : 2.23 (6H, s), 2.97 (2H, s), 4.79 (2H, s), 6.30 (1H, s), 6.96 (1H, d, J = 8.1 Hz), 7.13 (1H, s), 7.20 (1H, d, J = 8.1 Hz), 7.45 (2H, d, J = 8.6 Hz), 7.56 (2H, d, J = 8.6 Hz), 7.66 (2H, d, J = 8.4 Hz), 7.74 (1H, brs), 7.93 (2H, d, J = 8.4 Hz).

Melting point: 199 - 208°C (crystallization solvent: diisopropyl ether)

Example 91

15 2',4'-Difluoro-N-[3-[N,N-dimethylamino)methyl]-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide



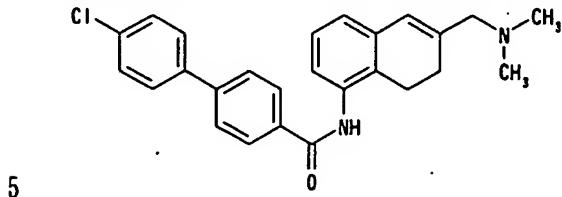
The titled compound was obtained by carrying out the same operation as in Example 1, using 3-[N,N-dimethylamino)methyl]-2H-chromen-7-amine obtained in Reference Example 59.

¹H-NMR (CDCl₃) δ : 2.23 (6H, s), 2.97 (2H, s), 4.78 (2H, s), 6.29 (1H, s), 6.80-7.10 (2H, m), 6.96 (1H, d, J = 8.1 Hz), 7.13 (1H, s), 7.20 (1H, d, J = 8.1 Hz), 7.40-7.50 (1H, m), 7.62 (2H, d, J = 8.4 Hz), 7.76 (1H, brs), 7.92 (2H, d, J = 8.4 Hz).

Melting point: 200 - 204°C (crystallization solvent: diisopropyl ether)

Example 92

4'-Chloro-N-[6-[(dimethylamino)methyl]-7,8-dihydro-1-naphthalenyl][1,1'-biphenyl]-4-carboxamide



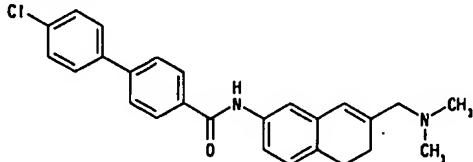
The titled compound was obtained in the same manner as in Example 1, using 6-[(dimethylamino)methyl]-7,8-dihydro-1-naphthalenamine obtained in Reference Example 10 60.

¹H-NMR (CDCl₃) δ : 2.34 (6H, s), 2.36 (2H, t, J=8.1 Hz), 2.80 (2H, t, J=8.1 Hz), 3.00 (2H, s), 6.38 (1H, s), 6.94 (1H, d, J=7.8 Hz), 7.21 (1H, t, J=7.8 Hz), 7.45 (2H, d, J=8.6 Hz), 7.56 (2H, d, J=8.6 Hz), 7.61 (2H, m), 7.68 (2H, d, J=8.4 Hz), 7.97 (2H, d, J=8.4 Hz).

Melting point: 193 - 195°C (crystallization solvent : diisopropyl ether)

Example 93

20 4'-Chloro-N-[7-[(dimethylamino)methyl]-5,6-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as a white powder by the same method as in Example 1, using 7-[(dimethylamino)methyl]-5,6-dihydro-2-naphthalenamine obtained in Reference Example 61.

¹H-NMR (CDCl₃) δ : 2.25 (6H, s), 2.34 (2H, t, J=8.1 Hz), 2.82 (2H, t, J=8.1 Hz), 3.00 (2H, s), 6.36 (1H, s), 7.11 (1H, d, J=7.5 Hz), 7.34 (1H, d, J=8.1 Hz), 7.38 (1H, s), 7.44

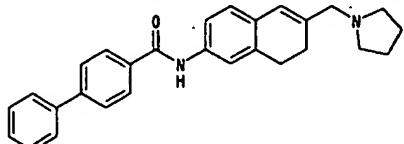
(2H, d, J=8.4 Hz), 7.56 (2H, d, J=8.4 Hz), 7.66 (2H, d, J=8.4 Hz), 7.78 (1H, brs), 7.97 (2H, d, J=8.4 Hz).

Melting point: 167 - 169°C (crystallization solvent : diisopropyl ether)

5

Example 94

N-[6-(1-Pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



10 The titled compound was obtained as a white powder in the same manner as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ : 1.75-1.90 (4H, m), 2.34 (2H, t, J=8.1 Hz), 2.45-2.60 (4H, m), 2.85 (2H, t, J=8.1 Hz), 3.18 (2H, s), 6.36 (1H, s), 7.02 (1H, d, J=8.1 Hz), 7.27-7.55 (5H, m), 7.63 (2H, d, J=7.3 Hz), 7.70 (2H, d, J=8.4 Hz), 7.82 (1H, s), 7.94 (2H, d, J=8.1 Hz).

Elemental analysis for C₂₈H₂₈N₂O

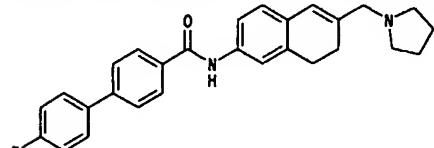
20 Calcd.: C, 82.32; H, 6.91; N, 6.86.

Found: C, 81.99; H, 6.69; N, 6.91.

Melting point: 176 - 177°C (crystallization solvent : diisopropyl ether)

25 **Example 95**

4'-Fluoro-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



30 The titled compound was obtained in the same manner as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-

dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ : 1.75-1.90 (4H, m), 2.35 (2H, t, J=8.2 Hz), 2.45-2.60 (4H, m), 2.84 (2H, t, J=8.2 Hz), 3.18 (2H, s), 5 6.36 (1H, s), 7.01 (1H, d, J=8.1 Hz), 7.16 (2H, t, J=8.1 Hz), 7.38 (1H, d, J=8.1 Hz), 7.48 (1H, brs), 7.56-7.61 (2H, m), 7.64 (2H, d, J=8.4 Hz), 7.83 (1H, s), 7.93 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₈H₂₇FN₂O

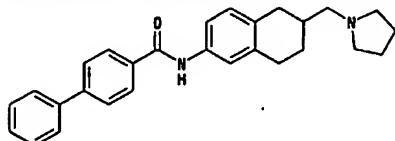
10 Calcd.: C, 78.85; H, 6.38; N, 6.57.

Found: C, 78.75; H, 6.39; N, 6.45.

Melting point: 189 - 192°C (crystallization solvent : diisopropyl ether)

15 Example 96

N-[6-(1-Pyrrolidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as a white powder in 20 the same manner as in Example 1, using 6-(1-pyrrolidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenamine obtained in Reference Example 55.

¹H-NMR (CDCl₃) δ : 1.40-1.50 (1H, m), 1.80 (4H, m), 1.80-2.10 (1H, m), 1.80-2.20 (8H, m), 3.30-4.00 (3H, m), 7.29 (1H, d, J=8.4 Hz), 7.25-7.30 (1H, m), 7.30-7.55 (4H, m), 6.43 (2H, d, J=7.0 Hz), 7.70 (2H, t, J=8.4 Hz), 7.75 (1H, s), 7.94 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₈H₃₀N₂O

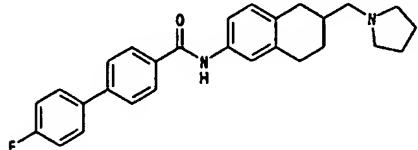
Calcd.: C, 81.91; H, 7.37; N, 6.82.

30 Found: C, 81.53; H, 7.25; N, 6.86.

Melting point: 144 - 146°C (crystallization solvent : diisopropyl ether)

Example 97

4'-Fluoro-N-[6-(1-pyrrolidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



5 The titled compound was obtained as a white powder in
the same manner as in Example 1, using 6-(1-
pyrrolidinylmethyl)-5,6,7,8-tetrahydro-2-
naphthalenamine obtained in Reference Example 55.

10 ¹H-NMR (CDCl₃) δ : 1.40-1.50 (1H, m), 1.80 (4H, m), 1.80-2.10
10 (1H, m), 1.80-2.20 (8H, m), 3.30-4.00 (3H, m), 7.08 (1H,
d, J=8.1 Hz), 7.15 (2H, t, J=8.4 Hz), 7.30 (1H, d, J=8.1
Hz), 7.44 (1H, brs), 7.56-7.61 (2H, m), 7.62 (2H, d, J=8.1
Hz), 7.85 (1H, s), 7.92 (2H, d, J=8.1 Hz).

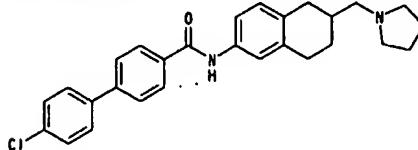
Elemental analysis for C₂₈H₂₉FN₂O

15 Calcd.: C, 78.48; H, 6.82; N, 6.54.
Found: C, 78.18; H, 6.60; N, 6.60.

Melting point: 185 - 189°C (crystallization solvent :
diisopropyl ether)

20 **Example 98**

4'-Chloro-N-[6-(1-pyrrolidinylmethyl)-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



25 The titled compound was obtained as a white powder in
the same manner as in Example 1, using 6-(1-
pyrrolidinylmethyl)-5,6,7,8-tetrahydro-2-
naphthalenamine obtained in Reference Example 55.

30 ¹H-NMR (CDCl₃) δ : 1.40-1.50 (1H, m), 1.80 (4H, m), 1.80-2.10
30 (1H, m), 1.80-2.20 (8H, m), 3.30-4.00 (3H, m), 7.08 (1H,
d, J=8.1 Hz), 7.31 (1H, d, J=8.4 Hz), 7.43 (2H, d, J=8.7
Hz), 7.45 (1H, s), 7.54 (2H, d, J=8.7 Hz), 7.64 (2H, d, J=8.4

Hz), 7.80 (1H, s), 7.93 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₈H₃₉ClN₂O

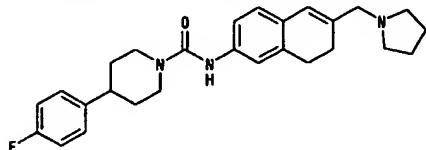
Calcd.: C, 75.57; H, 6.57; N, 6.30.

Found: C, 75.26; H, 6.68; N, 6.15.

5 Melting point: 206 - 209°C (crystallization solvent : diisopropyl ether)

Example 99

10 4-(4-Fluorophenyl)-N-[6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



15 6-(1-Pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54 (50 mg, 0.22 mmol) and pyridine (35 mg, 0.44 mmol) were dissolved in tetrahydrofuran (3 ml). Phenyl chlorocarbonate (38 mg, 0.24 mol) was added to the solution under ice-cooling, which was stirred for 10 minutes. The reaction mixture was concentrated, and dimethylsulfoxide (5 ml) was added to the residue. 4-(4-Fluorophenyl)piperidine hydrochloride (57 mg, 0.26 mmol) and 4N aqueous sodium hydroxide solution (0.066 ml, 0.26 mmol) were added to the reaction mixture at room temperature while stirring, which was stirred for 30 minutes. Ethyl acetate and water were added to the mixture, and extraction was conducted. The organic layer was washed with water, and concentrated. Diisopropyl ether was added to the residue. The crystallized product was collected by filtration, washed with diisopropyl ether, to give 4-(4-fluorophenyl)-N-[6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide (48 mg) as a white powder.

¹H-NMR (CDCl₃) δ: 1.60-1.70 (2H, m), 1.79 (4H, m), 1.80-1.90 (2H, m), 2.33 (2H, t, J=7.8 Hz), 2.51 (4H, m), 2.60-2.70 (1H, m), 2.80 (2H, t, J=7.8 Hz), 2.90-3.10 (2H, m), 3.16

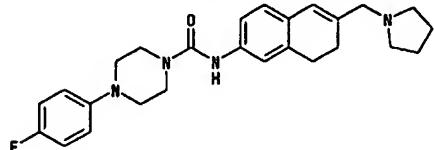
(2H, s), 4.18-4.23 (2H, m), 6.32 (1H, s), 6.32 (1H, s),
6.92-7.09 (4H, m), 7.15-7.20 (3H, m).

Melting point: 182 - 185°C (crystallization solvent :
diisopropyl ether)

5

Example 100

4-(4-Fluorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperazinecarboxamide



10 The titled compound was obtained as a white powder in the same manner as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54 and 4-fluorophenylpiperazine.

15 $^1\text{H-NMR}$ (CDCl_3) δ : 1.79 (4H, m), 2.32 (2H, t, $J=7.8$ Hz), 2.51(4H, m), 2.80 (2H, t, $J=7.8$ Hz), 3.13-3.16 (4H, m), 3.16 (2H, s), 3.63-3.66 (4H, m), 6.30 (1H, s), 6.32 (1H, s), 6.88-7.08 (6H, m), 7.19 (1H, s).

Elemental analysis for $\text{C}_{26}\text{H}_{31}\text{FN}_4\text{O}$

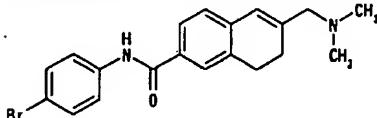
20 Calcd.: C, 71.86; H, 7.19; N, 12.89.

Found: C, 71.68; H, 7.35; N, 12.65.

Melting point: 179 - 181°C (crystallization solvent : diisopropyl ether)

25 **Example 101**

N-(4-Bromophenyl)-6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenecarboxamide



1) 6-Cyano-1-tetralone (1.30 g, 7.59 mmol)

30 synthesized by a known method by documents (synthetic communications, 23(21), 2965 (1993)) was dissolved in a

mixed solution of concentrated hydrochloric acid (10 ml) and acetic acid (20 ml), which was stirred at 120°C for 16 hours. The reaction mixture was concentrated. Ethyl acetate and water were added to the residue, and extraction 5 was conducted. The organic layer was washed with water, and concentrated. The residue was washed with ethyl acetate - n-hexane (1:1), to give 5-oxo-5,6,7,8-tetrahydro-2-naphthalenecarboxylic acid (1.10 g) as a white powder.

10 $^1\text{H-NMR}$ (CDCl_3) δ : 2.15-2.23 (2H, m), 2.70-2.75 (2H, m), 3.04-3.07 (2H, m), 8.01-8.03 (1H, m), 8.03 (1H, s), 8.13 (1H, d, $J=8.7$ Hz).

15 2) $\text{N-(4-Bromophenyl)-5-oxo-5,6,7,8-tetrahydro-2-naphthalenecarboxamide}$ (1.21 g) was obtained as a white powder in the same manner as in Example 1, using 5-oxo-5,6,7,8-tetrahydro-2-naphthalenecarboxylic acid (1.00 g, 5.26 mmol) obtained in 1) and 4-bromoaniline (0.90 g, 5.26 mmol).

20 $^1\text{H-NMR}$ (CDCl_3) δ : 2.14-2.23 (2H, m), 2.69-2.73 (2H, m), 3.03-3.07 (2H, m), 7.48-7.58 (4H, m), 7.71 (1H, d, $J=8.1$ Hz), 7.79 (1H, s), 7.86 (1H, s), 8.12 (1H, d, $J=8.1$ Hz).

25 3) $\text{N-(4-Bromophenyl)-5-oxo-5,6,7,8-tetrahydro-2-naphthalenecarboxamide}$ (1.10 g, 3.19 mmol) obtained in 2) was dissolved in dimethylformamide diethylacetal (30 ml), which was refluxed with heating for 4 hours. The crystallized product was collected by filtration, washed with ethyl acetate, to give $\text{N-(4-bromophenyl)-6-[(dimethylamino)methylidene]-5-oxo-5,6,7,8-tetrahydro-2-naphthalenecarboxamide}$ (1.21 g) as a yellow powder.

30 $^1\text{H-NMR}$ (CDCl_3) δ : 2.80-2.87 (4H, m), 3.07 (6H, m), 7.46-7.72 (7H, m), 7.91 (1H, d, $J=8.4$ Hz), 8.53 (1H, s).

35 4) Sodium triacetoxyhydroborate (398 mg, 1.87 mmol) was dissolved in a mixed solution of acetic acid (40 ml) and tetrahydrofuran (10 ml) under ice-cooling. $\text{N-(4-Bromophenyl)-6-[(dimethylamino)methylidene]-5-oxo-5,6,7,8-tetrahydro-2-naphthalenecarboxamide}$ (500 mg,

1.25 mmol) obtained in 3) was added to the solution, which was stirred for 1 hour. The reaction mixture was concentrated under reduced pressure at room temperature.

2-Propanol (50 ml) was added to the residue, and sodium borohydride (142 mg, 3.75 mmol) was further added under ice-cooling. After stirring for 2 hours, the reaction mixture was concentrated. Sodium hydrogencarbonate solution and ethyl acetate was added to the residue for liquid separation. The organic layer was concentrated.

10 The residue was dissolved in a mixed solution of acetic acid (20 ml) and concentrated hydrochloric acid (20 ml), which was stirred at 70°C for 5 hours. The reaction mixture was concentrated. 4N aqueous sodium hydroxide solution and ethyl acetate were added to the residue, and extraction was conducted. The organic layer was washed with water, and concentrated. The residue was purified by alumina column chromatography (development solvent: ethyl acetate), and the eluent was washed with diisopropyl ether, to give the titled compound (234 mg) as a white powder.

20 $^1\text{H-NMR}$ (CDCl_3) δ : 2.26 (6H, s), 2.38 (2H, t, $J=8.1$ Hz), 2.89 (2H, t, $J=8.1$ Hz), 3.02 (2H, s), 6.42 (1H, s), 7.10 (1H, d, $J=8.6$ Hz), 7.47 (2H, d, $J=8.9$ Hz), 7.55 (2H, d $J=8.9$ Hz), 7.61 (1H, s), 7.62 (1H, d, $J=6.7$ Hz), 7.76 (1H, s).

Elemental analysis for $\text{C}_{20}\text{H}_{21}\text{BrN}_2\text{O}$

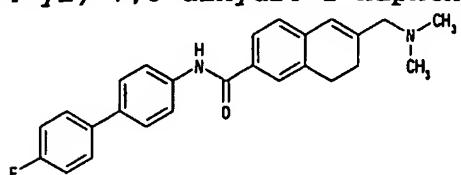
25 Calcd.: C, 62.35; H, 5.49; N, 7.27.

Found: C, 61.98; H, 5.43; N, 7.07.

Melting point: 175 - 179°C (crystallization solvent : diisopropyl ether)

30 Example 102

6-[(Dimethylamino)methyl]-N-(4'-fluoro[1,1'-biphenyl]-4-yl)-7,8-dihydro-2-naphthalenecarboxamide



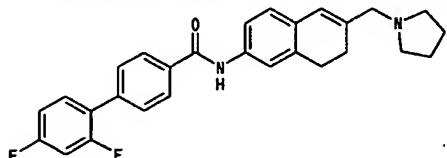
The titled compound was obtained as a white powder, by the same method as in Example 16, using N-(4-bromophenyl)-6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenecarboxamide (170 mg, 0.44 mmol) obtained in Example 101 and 4-fluorophenylboric acid (74 mg, 0.53 mmol).

$^1\text{H-NMR}$ (CDCl_3) δ : 2.27 (6H, s), 2.39 (2H, t, $J=8.4$ Hz), 2.91 (2H, t, $J=8.4$ Hz), 3.02 (2H, s), 6.43 (1H, s), 7.09-7.16 (3H, m), 7.52-7.73 (8H, m), 7.81 (1H, s).

Melting point: 200 - 204°C (crystallization solvent : diisopropyl ether)

Example 103

2',4'-Difluoro-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as a white powder by the same method as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

$^1\text{H-NMR}$ (CDCl_3) δ : 1.75-1.90 (4H, m), 2.36 (2H, t, $J=8.1$ Hz), 2.45-2.60 (4H, m), 2.85 (2H, t, $J=8.1$ Hz), 3.18 (2H, s), 6.36 (1H, s), 6.92-7.03 (3H, m), 7.36-7.45 (2H, m), 7.48 (1H, s), 7.62 (2H, d, $J=8.4$ Hz), 7.78 (1H, s), 7.94 (2H, d, $J=8.4$ Hz).

Elemental analysis for $\text{C}_{28}\text{H}_{26}\text{F}_2\text{N}_2\text{O}$

Calcd.: C, 75.66; H, 5.90; N, 6.30.

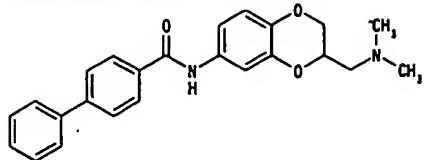
Found: C, 75.36; H, 5.92; N, 6.10.

Melting point: 165 - 167°C (crystallization solvent : diisopropyl ether)

Example 104

N-[3-[(Dimethylamino)methyl]-2,3-dihydro-1,4-

benzodioxin-6-yl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as a white powder by the same method as in Example 1, using N,N-dimethyl-N-
5 [(7-amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine obtained in Reference Example 62.

¹H-NMR (CDCl₃) δ : 2.33 (6H, s), 2.48-2.66 (2H, m), 3.93-3.99
10 (1H, m), 4.27-4.31 (2H, m), 6.86 (1H, d, J=8.6 Hz), 7.03-7.07
(1H, m), 7.31-7.32 (1H, m), 7.37-7.49 (3H, m), 7.62 (2H,
d, J=7.0 Hz), 7.68 (2H, d, J=8.4 Hz), 7.76 (1H, s), 7.91 (2H,
d, J=8.4 Hz).

Elemental analysis for C₂₄H₂₄N₂O₃

Calcd. : C, 74.21; H, 6.23; N, 7.21.

Found : C, 74.17; H, 6.23; N, 7.01.

15 Melting point: 124 - 126°C (crystallization solvent : diisopropyl ether)

Example 105

4'-Chloro-N-[3-[(dimethylamino)methyl]-2,3-dihydro-1,4-
20 benzodioxin-6-yl][1,1'-biphenyl]-4-carboxamide

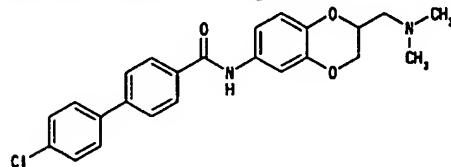
The titled compound was obtained as a white powder by the same method as in Example 1, using N,N-dimethyl-N-
25 [(7-amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine obtained in Reference Example 62.

¹H-NMR (CDCl₃) δ : 2.33 (6H, s), 2.50-2.67 (2H, m), 3.94-4.01
10 (1H, m), 4.28-4.31 (2H, m), 6.86 (1H, d, J=8.7 Hz),
7.03-7.06 (1H, m), 7.31 (1H, m), 7.44 (2H, d, J=8.4 Hz),
7.55 (2H, d, J=8.4 Hz), 7.65 (2H, d, J=8.1 Hz), 7.67 (1H,
30 s), 7.91 (2H, d, J=8.1 Hz).

Melting point: 158 - 159°C (crystallization solvent : diisopropyl ether)

Example 106

5 4'-Chloro-N-[2-[(dimethylamino)methyl]-2,3-dihydro-1,4-benzodioxin-6-yl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as a white powder by the same method as in Example 1, using N,N-dimethyl-N-
10 [(6-amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine obtained in Reference Example 63.

¹H-NMR (CDCl₃) δ : 2.34 (6H, s), 2.46-2.67 (2H, m), 3.94-4.01
15 (1H, m), 4.28-4.34 (2H, m), 6.91 (1H, d, J=8.6 Hz),
7.02-7.05 (1H, m), 7.30 (1H, m), 7.44 (2H, d, J=8.4 Hz),
7.55 (2H, d, J=8.4 Hz), 7.66 (2H, d, J=8.1 Hz), 7.70 (1H,
s), 7.92 (2H, d, J=8.1 Hz).

Elemental analysis for C₂₄H₂₃ClN₂O₃

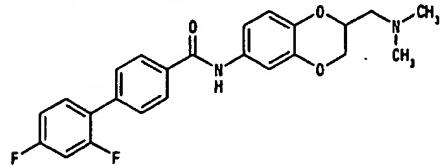
Calcd.: C, 68.16; H, 5.48; N, 6.62.

Found: C, 68.09; H, 5.29; N, 6.57.

20 Melting point: 215 - 217°C (crystallization solvent : diisopropyl ether)

Example 107

2',4'-Difluoro-N-[2-[(dimethylamino)methyl]-2,3-dihydro-1,4-benzodioxin-6-yl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as a white powder by the same method as in Example 1, using N,N-dimethyl-N-
30 [(6-amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]amine

obtained in Reference Example 63.

¹H-NMR (CDCl₃) δ : 2.34 (6H, s), 2.50-2.63 (2H, m), 3.94-4.01 (1H, m), 4.28-4.34 (2H, m), 6.91 (1H, d, J=8.6 Hz), 6.91-7.03 (3H, m), 7.30 (1H, m), 7.40-7.50 (1H, m), 7.61 (2H, d, J=8.1 Hz), 7.69 (1H, s), 7.92 (2H, d, J=8.1 Hz).

Elemental analysis for C₂₄H₂₂F₂N₂O₃

Calcd. : C, 67.91; H, 5.22; N, 6.60.

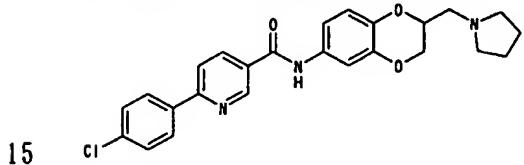
Found : C, 67.75; H, 5.09; N, 6.48.

Melting point: 209 - 210°C (crystallization solvent :

10 diisopropyl ether)

Example 108

6-(4-Chlorophenyl)-N-[2-(1-pyrrolidinylmethyl)-2,3-dihydro-1,4-benzodioxin-6-yl]nicotinamide



15

The titled compound was obtained as a white powder by the same method as in Example 1, using 1-[(6-amino-2,3-dihydro-1,4-benzodioxin-2-yl)methyl]pyrrolidine obtained in Reference Example 64.

20

¹H-NMR (CDCl₃) δ : 1.81 (4H, m), 2.50-2.63 (4H, m), 2.75-2.77 (2H, m), 3.90-4.10 (1H, m), 4.30-4.36 (2H, m), 6.91 (1H, d, J=8.6 Hz), 7.00-7.10 (1H, m), 7.26 (1H, m), 7.48 (2H, d, J=8.6 Hz), 7.72 (1H, s), 7.81 (1H, d, J=7.8 Hz), 8.01 (2H, d, J=8.6 Hz), 8.20-8.25 (1H, m), 9.10 (1H, s).

25

Elemental analysis for C₂₅H₂₄ClN₃O₃

Calcd. : C, 66.74; H, 5.38; N, 9.34.

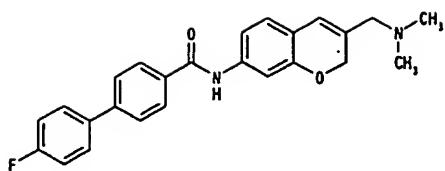
Found : C, 66.66; H, 5.46; N, 9.11.

Melting point: 218 - 220°C (crystallization solvent : diisopropyl ether)

30

Example 109

N-[3-[(Dimethylamino)methyl]-2H-chromen-7-yl]-4'-fluoro[1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 3-[(N,N-dimethylamino)methyl]-2H-chromen-7-amine obtained in

5 Reference Example 59.

¹H-NMR (CDCl₃) δ : 2.23 (6H, s), 2.97 (2H, s), 4.79 (2H, s), 6.30 (1H, s), 6.96 (1H, d, J=8.1 Hz), 7.13-7.22 (4H, m), 7.56-7.61 (2H, m), 7.65 (2H, d, J=8.4 Hz), 7.78 (1H, s), 7.92 (2H, d, J=8.4 Hz).

10 Elemental analysis for C₂₅H₂₃FN₂O₂

Calcd.: C, 74.61; H, 5.76; N, 6.96.

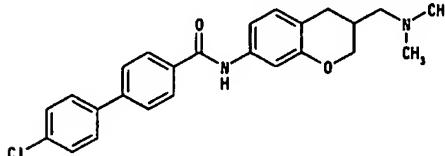
Found: C, 74.35; H, 5.68; N, 6.74.

Melting point: 192 - 195°C (crystallization solvent : diisopropyl ether)

15

Example 110

4'-Chloro-N-[3-[(dimethylamino)methyl]-3,4-dihydro-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide

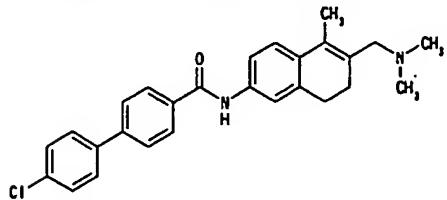


20 The titled compound was obtained by carrying out the same operation as in Example 1, using N-[(7-amino-3,4-dihydro-2H-chromen-3-yl)methyl]-N,N-dimethylamine obtained in Reference Example 65.

¹H-NMR (CDCl₃) δ : 2.26 (6H, s), 2.27 (3H, m), 2.47-2.51 (1H, m), 2.83-2.89 (1H, m), 3.82-3.86 (1H, m), 4.28-4.32 (1H, m), 7.04 (1H, d, J=8.1 Hz), 7.12-7.18 (2H, m), 7.44 (2H, d, J=8.4 Hz), 7.56 (2H, d, J=8.4 Hz), 7.67 (2H, d, J=8.4 Hz), 7.71 (1H, s), 7.93 (2H, d, J=8.4 Hz).

30 Example 111

4'-Chloro-N-[6-[(dimethylamino)methyl]-5-methyl-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the
 5 same operation as in Example 1, using 6-
 [(dimethylamino)methyl]-5-methyl-7,8-dihydro-2-
 naphthalenamine obtained in Reference Example 66.

¹H-NMR (CDCl₃) δ : 2.09 (3H, s), 2.27 (6H, s), 2.31-2.37 (2H, m), 2.74-2.79 (2H, m), 3.08 (2H, s), 7.27-7.30 (1H, m),
 10 7.44-7.48 (4H, m), 7.56 (2H, d, J=8.6 Hz), 7.67 (2H, d, J=8.4 Hz), 7.79 (1H, s), 7.95 (2H, d, J=8.4 Hz).

Elemental analysis for C₂₇H₂₇ClN₂O

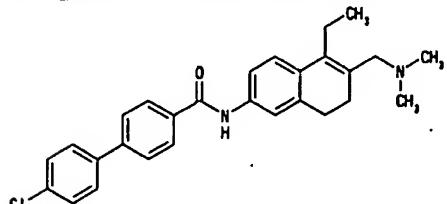
Calcd. : C, 75.25; H, 6.31; N, 6.50.

Found : C, 74.86; H, 6.20; N, 6.42.

15 Melting point: 199 - 204°C (crystallization solvent :
 diisopropyl ether)

Example 112

4'-Chloro-N-[6-[(dimethylamino)methyl]-5-ethyl-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the
 same operation as in Example 1, using 6-
 [(dimethylamino)methyl]-5-ethyl-7,8-dihydro-2-
 naphthalenamine obtained in Reference Example 67.

¹H-NMR (CDCl₃) δ : 1.09 (3H, t, J=7.5 Hz), 2.27 (6H, s),
 2.31-2.37 (2H, m), 2.60-2.63 (2H, m), 2.71-2.76 (2H, m),
 3.08 (2H, s), 7.31 (1H, d, J=9.2 Hz), 7.43-7.49 (4H, m),

7.56 (2H, d, J=8.7 Hz), 7.67 (2H, d, J=8.6 Hz), 7.80 (1H, s), 7.94 (2H, d, J=8.6 Hz).

Elemental analysis for C₂₈H₂₉ClN₂O

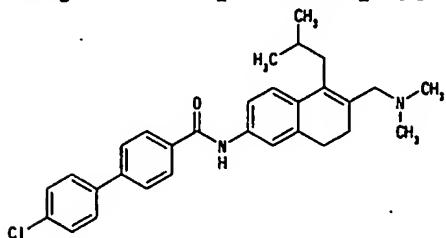
Calcd.: C, 75.57; H, 6.57; N, 6.30.

5 Found: C, 75.41; H, 6.34; N, 6.23.

Melting point: 201 - 204°C (crystallization solvent : diisopropyl ether)

Example 113

10 4'-Chloro-N-[6-[(dimethylamino)methyl]-5-isobutyl-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-

15 [(dimethylamino)methyl]-5-isobutyl-7,8-dihydro-2-naphthalenamine obtained in Reference Example 68.

1H-NMR (CDCl₃) δ: 0.90 (6H, d, J=6.4 Hz), 1.73-1.78 (1H, m), 2.23 (6H, s), 2.34 (2H, m), 2.50 (2H, d, J=7.3 Hz), 2.74 (2H, m), 3.13 (2H, s), 7.26-7.30 (1H, m), 7.45-7.48 (4H, m), 7.56 (2H, d, J=8.7 Hz), 7.67 (2H, d, J=8.4 Hz), 7.79 (1H, s), 7.94 (2H, d, J=8.4 Hz).

Elemental analysis for C₃₀H₃₃ClN₂O

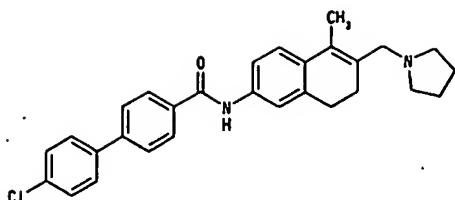
Calcd.: C, 76.17; H, 7.03; N, 5.92.

Found: C, 75.91; H, 7.19; N, 5.72.

25 Melting point: 159 - 162°C (crystallization solvent : diisopropyl ether)

Example 114

4'-Chloro-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 5-methyl-6- (1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine

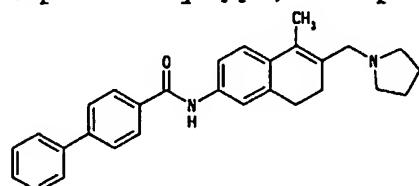
5 obtained in Reference Example 69.

¹H-NMR (CDCl₃) δ : 1.79 (4H, m), 2.11 (3H, s), 2.30-2.40 (2H, m), 2.54 (4H, m), 2.74-2.79 (2H, m), 3.28 (2H, s), 7.26-7.30 (1H, m), 7.45-7.48 (4H, m), 7.56 (2H, d, J=8.6 Hz), 7.67 (2H, d, J=8.4 Hz), 7.81 (1H, s), 7.95 (2H, d, J=8.4 Hz).

Melting point: 190 - 192°C (crystallization solvent : diisopropyl ether)

Example 115

15 N-[5-Methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



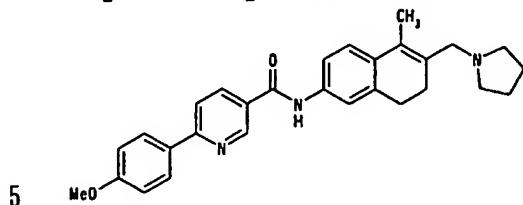
The titled compound was obtained by carrying out the same operation as in Example 1, using 5-methyl-6- (1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine
20 obtained in Reference Example 69.

¹H-NMR (CDCl₃) δ : 1.78 (4H, m), 2.10 (3H, s), 2.35-2.40 (2H, m), 2.53 (4H, m), 2.70-2.78 (2H, m), 3.28 (2H, s), 7.26-7.28 (1H, m), 7.40-7.50 (5H, m), 7.62 (2H, d, J=7.0 Hz), 7.70 (2H, d, J=8.4 Hz), 7.87 (1H, s), 7.94 (2H, d, J=8.4 Hz).

Melting point: 169 - 170°C (crystallization solvent : diisopropyl ether)

Example 116

6-(4-Methoxyphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

10 ¹H-NMR (CDCl₃) δ : 1.78 (4H, m), 2.09 (3H, s), 2.35-2.40 (2H, m), 2.53 (4H, m), 2.70-2.77 (2H, m), 3.27 (2H, s), 3.88 (3H, s), 7.01 (2H, d, J=8.9 Hz), 7.26 (1H, d, J=8.9 Hz), 7.45-7.47 (2H, m), 7.75 (1H, d, J=8.4 Hz), 7.95 (1H, s), 8.01 (2H, d, J=8.9 Hz), 8.18-8.21 (1H, m), 9.09 (1H, m).

15 Elemental analysis for C₂₉H₃₁N₃O₂

Calcd. : C, 76.79; H, 6.89; N, 9.26.

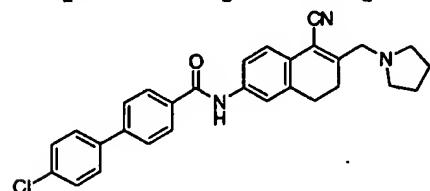
Found : C, 76.46; H, 6.64; N, 9.09.

Melting point: 165 - 167°C (crystallization solvent : diisopropyl ether)

20

Example 117

4'-Chloro-N-[5-cyano-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



25 The titled compound was obtained as a colorless powder by carrying out the same operation as in Example 1, using 6-amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-1-naphthalenecarbonitrile obtained in Reference Example 70 and 4'-chloro[1,1'-biphenyl]-4-carboxylic acid.

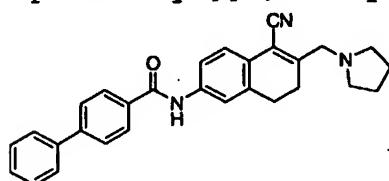
¹H NMR (DMSO-d₆) δ : 1.73 (4H, m), 2.50 (4H, m), 2.56 (2H, m), 2.82 (2H, m), 3.49 (2H, s), 7.32 (1H, d, J = 9.0 Hz), 7.57 (2H, d, J = 8.4 Hz), 7.56-7.87 (6H, m), 8.07 (2H, d, J = 8.4 Hz), 10.40 (1H, s).

5 FABMS(pos) 468.2 [M+H]⁺

Melting point: 191 - 192°C (crystallization solvent : diisopropyl ether)

Example 118

10 N-[5-Cyano-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, 15 using 6-amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-1-naphthalenecarbonitrile obtained in Reference Example 70 and [1,1'-biphenyl]-4-carboxylic acid.

1 H NMR (DMSO-d₆) δ : 1.81 (4H, m), 2.62 (6H, m), 2.88 (2H, m), 3.56 (2H, s), 7.41 (2H, m), 7.46 (3H, m), 7.64 (2H, d, 20 J = 6.9 Hz), 7.73 (3H, m), 7.88 (1H, s), 7.95 (2H, d, J = 8.1 Hz).

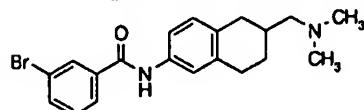
FABMS(pos) 434.2 [M+H]⁺

Melting point: 168 - 170°C (crystallization solvent : diisopropyl ether)

25

Example 119

3-Bromo-N-[6-[(dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide

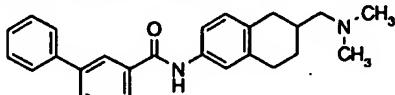


30 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-amino-2-[(N,N-

dimethylamino)methyl]tetralin and 3-bromobenzoic acid.
¹H NMR (DMSO-d₆) δ : 1.31 (1H, m), 1.89 (2H, m), 2.17 (6H, s), 2.17-2.35 (3H, m), 2.77 (3H, m), 7.04 (1H, d, J=8.4 Hz), 7.49 (3H, m), 7.79 (1H, d, J=8.1 Hz), 7.94 (1H, d, J=7.8 Hz), 8.13 (1H, s), 10.20 (1H, s).

Example 120

N-[6-[(Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-3-carboxamide



10

The titled compound was obtained by carrying out the same operation as in Example 16, using 3-bromo-N-[6-[(dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]benzamide obtained in Example 119 and phenylboronic acid.

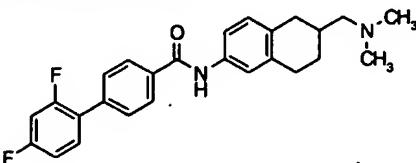
¹H NMR (DMSO-d₆) δ : 1.43 (1H, m), 2.02 (1H, m), 2.21 (1H, m), 2.42 (1H, m), 2.81 (6H, s), 2.88 (3H, m), 3.09 (2H, m), 7.06 (1H, m), 7.42-7.65 (6H, m), 7.78-7.95 (4H, m), 8.22 (1H, s), 10.27 (1H, s).

20 FABMS(pos) 385.2 [M+H]⁺

Melting point: 145 - 148°C (crystallization solvent : ethyl acetate-diisopropyl ether)

Example 121

25 N-[6-[(Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-2',4'-difluoro[1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin and 2', 4'-difluoro[1,1'-

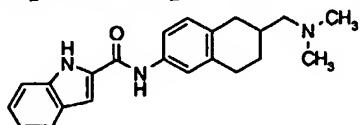
biphenyl]-4-carboxylic acid.

¹H NMR (CDCl₃) δ : 1.41 (1H, m), 1.94 (2H, m), 2.25 (6H, s), 2.23-2.30 (3H, m), 2.86 (3H, m), 6.96 (2H, m), 7.09 (1H, d, J=8.1 Hz), 7.30 (1H, m), 7.43 (2H, m), 7.61 (2H, m), 7.76
5 (1H, s), 7.93 (2H, m).

Melting point: 162 - 163°C (crystallization solvent : ethyl acetate-diisopropyl ether)

Example 122

10 N-[6-[(Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl-1H-indole-2-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin and 1H-indol-2-carboxylic acid.

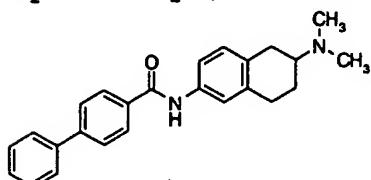
¹H NMR (DMSO-d₆) δ : 1.32 (1H, m), 1.91 (2H, m), 2.16 (6H, s), 2.16-2.35 (3H, m), 2.78 (3H, m), 7.06 (2H, m), 7.21 (1H, m), 7.44 (4H, m), 7.66 (1H, d, J=8.1 Hz), 10.05 (1H, s),
20 11.68 (1H, s).

FABMS(pos) 348.2 [M+H]⁺

Melting point: 190 - 192°C (crystallization solvent : ethyl acetate - diisopropyl ether)

25 Example 123

N-[6-[(Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl] [1,1'-biphenyl]-4-carboxamide

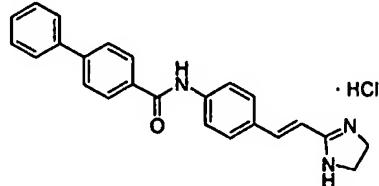


A tetrahydrofuran solution (0.146ml, 0.293mmol) of
30 N-(6-oxo-5,6,7,8-tetrahydro-2-naphthalenyl)[1,1'-

biphenyl]-4-carboxamide (10 mg, 0.029 mmol) obtained in Reference Example 72 and 2N dimethylamine was added to acetic acid-tetrahydrofuran (1:1) solution (0.5ml), which was stirred at 50°C for 15 minutes. After the reaction 5 mixture was cooled at room temperature, sodium triacetoxyhydroborate (31 mg, 0.146 mmol) was added, which was stirred at 50°C for 2 hours. 1N Hydrochloric acid was added to the reaction mixture, which was washed with ethyl acetate. Sodium carbonate was added to the water layer to 10 make it alkaline, then extraction was conducted using ethyl acetate. The extract was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was purified by 15 alumina B column chromatography (development solvent; ethyl acetate), to give the titled compound (1.6mg).
¹H NMR (CDCl₃) δ: 1.68 (1H, m), 2.27 (1H, m), 2.40 (6H, s), 2.78 (5H, m), 7.11 (1H, d, J=8.1 Hz), 7.32-7.50 (5H, m), 7.62 (2H, m), 7.72 (2H, d, J=8.4 Hz), 7.78 (1H, br), 7.94 (2H, d, J=8.4 Hz).
 20 FABMS(pos) 371.2 [M+H]⁺

Example 124

25 N-[4-[(E)-2-(4,5-Dihydro-1H-imidazol-2-yl)ethenyl]phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



10.1 N Hydrogen chloride-ethanol solution (30 ml) was added to an ethanol suspension of N-[4-[(E)-2-30 cyanoethenyl]phenyl][1,1'-biphenyl]-4-carboxamide (250 mg, 0.771 mmol) obtained in Reference Example under room temperature, which was stirred for 16 hours. After the

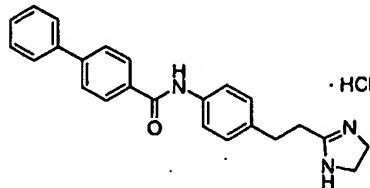
solvent was distilled out under reduced pressure, ethanol was again added to the residue, and then ethylenediamine (0.155 ml, 2.31 mmol) was added at room temperature, which was stirred for 16 hours. Sodium hydrogencarbonate solution was added to the reaction mixture, and the precipitated crude product was washed with water and chloroform. This product was dissolved in methanol. 1 N Hydrochloric acid (4 ml) was added to the solution, and the solvent was distilled out under reduced pressure.

Small amount of water was added to the resulting residue, to give the titled compound (124 mg) as a colorless powder.
¹H NMR (DMSO-d₆, free base) δ : 3.33 (4H, m), 6.61 (1H, d, J = 16.8 Hz), 7.15 (1H, d, J = 16.8 Hz), 7.52 (5H, m), 7.83 (6H, m), 8.07 (2H, d, J = 8.4 Hz).

Elemental analysis for C₂₄H₂₁N₃O · HCl · 1.5H₂O
 Calcd. : C, 66.89; H, 5.85; N, 9.75.
 Found : C, 67.16; H, 6.10; N, 10.03.

Example 125

N-[4-[(E)-2-(4,5-Dihydro-1H-imidazol-2-yl)ethenyl]phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



10% Palladium – carbon (200 mg) was added to a methanol suspension of N-[4-[(E)-2-(4,5-dihydro-1H-imidazol-2-yl)ethenyl]phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride (80 mg, 0.198 mmol) obtained in Example 124, which was stirred under hydrogen atmosphere at 60°C for 2 hours. After a catalyst was filtered off, the solvent was distilled out under reduced pressure. Diethyl ether was added to the resulting residue, to give the titled compound (52 mg) as a colorless powder.

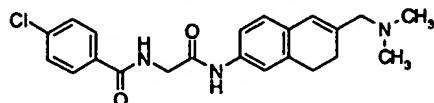
¹H NMR (DMSO-d₆) δ : 2.73-2.97 (4H, m), 3.37 (4H, s), 7.24 (2H, d, J = 8.4 Hz), 7.46 (3H, m), 7.76 (6H, m), 8.08 (2H, d, J = 8.4 Hz).

FABMS(pos) 370 [M+H]⁺

5

Example 126

4-Chloro-N-[2-[[6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]amino]-2-oxoethyl]benzamide



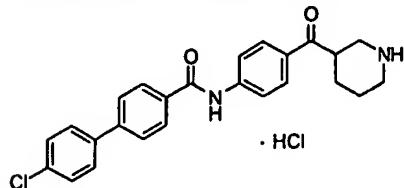
10 The titled compound was obtained by carrying out the same operation as in Example 1, using 6- [(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Example 41-2) and 4-chlorobenzoyl glycine.
¹H NMR (DMSO-d₆) δ : 2.18 (6H, s), 2.21 (2H, m), 2.71 (2H, m), 2.91 (2H, s), 4.05 (2H, d, J=5.6 Hz), 6.30 (1H, s), 6.98 (1H, d, J=8.1 Hz), 7.36 (2H, m), 7.58 (2H, d, J=8.4 Hz), 7.92 (2H, d, J=8.4 Hz), 8.94 (1H, t, J=5.6 Hz), 10.00 (1H, s).

FABMS(pos) 398 [M+H]⁺

15 Melting point: 168 - 171°C (crystallization solvent : diisopropyl ether)

Example 127

4'-Chloro-N-[4-(3-piperidinylcarbonyl)phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



20 1) tert-Butyl 3-[[4-[(4'-chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]benzoyl]-1-piperidinecarboxylate was obtained by carrying out the same operation as in Example 1, using tert-butyl 3-(4-aminobenzoyl)-1-piperidinecarboxylate obtained in Reference Example 77 and

4'-chloro[1,1'-biphenyl]-4-carboxylic acid.

FABMS(pos) 519.2 [M+H]⁺

2) 4N Hydrogen chloride-ethyl acetate (1 ml) was added to tert-butyl 3-[4-[(4'-chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]benzoyl]-1-piperidinecarboxylate (100 mg, 0.193 mmol) obtained in 1). One hour later, the solvent was distilled out under reduced pressure. Diisopropyl ether was added to the residue, to give the titled compound (73.3 mg) as a colorless powder.

10 ¹H NMR (DMSO-d₆) δ : 1.56 (1H, m), 1.82 (2H, m), 2.02 (1H, m), 2.89 (1H, m), 3.05 (1H, m), 3.33 (2H, m), 3.90 (1H, m), 7.58 (2H, d, J=8.1Hz), 7.81 (2H, d, J=8.1Hz), 7.88 (2H, d, J=8.1Hz), 8.03 (4H, m), 8.11 (2H, d, J=8.1Hz), 9.04 (2H, br), 10.73 (1H, s).

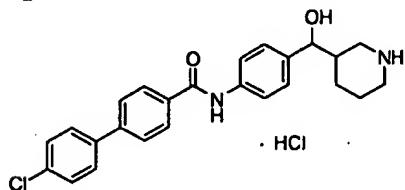
15 FABMS(pos) 419.2 [M+H]⁺

Melting point: 222 - 225°C (decomposition)

Example 128

4'-Chloro-N-[4-[hydroxy(3-

20 piperidinyl)methyl]phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



4N Hydrogen chloride-ethyl acetate (1 ml) was added to tert-butyl 3-[4-[(4'-chloro[1,1'-biphenyl]-4-

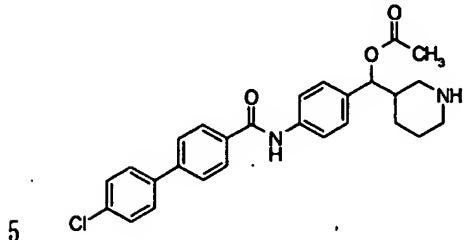
25 yl)carbonyl]amino]phenyl](hydroxy)methyl]-1-piperidinecarboxylate (100 mg, 0.192 mmol) obtained in Reference Example 78. One hour later, the solvent was distilled out under reduced pressure. Diisopropyl ether was added to the residue, to give the titled compound (79.8 mg) as a colorless powder.

30 FABMSMS(pos) 421.2 [M+H]⁺

Melting point: 195°C (decomposition)

Example 129

[4-[[[4'-Chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]phenyl](3-piperidinyl)methyl acetate



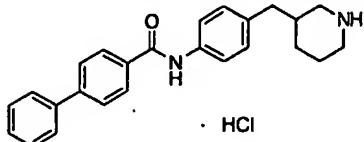
Concentrated sulfuric acid (0.0562 ml) was added to an acetic acid solution (3.5 ml) of tert-butyl 3-[[4-
5 [[(4'-chloro[1,1'-biphenyl]-4-
yl)carbonyl]amino]phenyl](hydroxy)methyl]-1-
10 piperidinecarboxylate (366 mg, 0.702 mmol) obtained in
Example 128, which was stirred under room temperature for
16 hours. Ethyl acetate was added to the reaction mixture,
which was washed with potassium hydrogencarbonate solution
and saturated aqueous sodium chloride solution, dried over
15 anhydrous sodium sulfate, and then the solvent was
distilled out under reduced pressure. The resulting oily
substance was purified by alumina B column chromatography
(development solvent; ethyl acetate : methanol = 3:1), and
powdered with diisopropyl ether, to give the titled
15 compound (210 mg).

20 FABMS(pos) 403.2 [M+H]⁺

Melting point: 200 - 203°C.

Example 130

25 N-[4-(3-Piperidinylmethyl)phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride

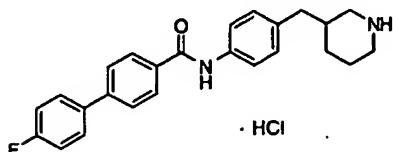


4N Hydrogen chloride-ethyl acetate (2 ml) was added to tert-butyl 3-[[4-[[1,1'-biphenyl]-4-

ylcarbonyl)amino]benzyl]-1-piperidinecarboxylate (100 mg, 0.212 mmol) obtained in Reference Example 80. Two hours later, the solvent was distilled out under reduced pressure. Diisopropyl ether was added to the residue for 5 powdering, to give the titled compound (79 mg).
 FABMS(pos) 371.3 [M+H]⁺
 Melting point: 218 - 220°C (decomposition)

Example 131

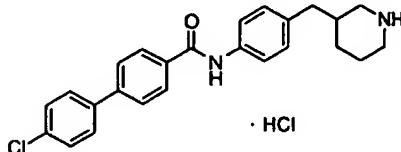
10 4'-Fluoro-N-[4-(3-piperidinylmethyl)phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



15 4 N Hydrogen chloride-ethyl acetate (3 ml) was added to tert-butyl 3-[4-[(4'-fluoro[1,1'-biphenyl]-4-yl)carbonyl]amino]benzyl]-1-piperidinecarboxylate (150 mg, 0.307 mmol) obtained in Reference Example 81. Two hours later, the solvent was distilled out under reduced pressure. Diisopropyl ether was added to the residue, to give the titled compound (115 mg) as a colorless powder.
 20 FABMS(pos) 389.3 [M+H]⁺
 Melting point: 205°C (decomposition)

Example 132

25 4'-Chloro-N-[4-(3-piperidinylmethyl)phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



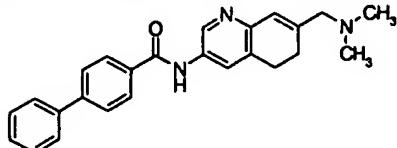
30 4 N Hydrogen chloride-ethyl acetate (3 ml) was added to tert-butyl 3-[4-[(4'-chloro[1,1'-biphenyl]-4-yl)carbonyl]amino]benzyl]-1-piperidinecarboxylate (150 mg, 0.297 mmol) obtained in Reference Example 82. Two hours

later, the solvent was distilled out under reduced pressure. Diisopropyl ether was added to the residue, to give the titled compound (73.3 mg) as a colorless powder.
 FABMS(pos) 405.2 [M+H]⁺

5 Melting point: 200°C (decomposition)

Example 133

N-[7-[(Dimethylamino)methyl]-5,6-dihydro-3-quinolinyl][1,1'-biphenyl]-4-carboxamide



10

The titled compound was obtained by carrying out the same operation as in Example 1, using N-[(3-amino-5,6-dihydro-7-quinolinyl)methyl]-N,N-dimethylamine obtained in Reference Example 86 and [1,1'-biphenyl]-4-carboxylic acid.

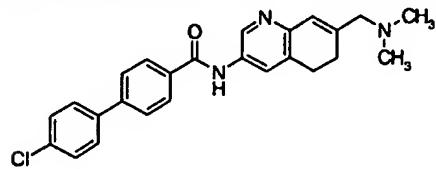
15 ¹H NMR (DMSO-d₆) δ : 2.16 (6H, s), 2.29 (2H, t, J=8.1 Hz), 2.84 (2H, t, J=8.1 Hz), 2.98 (2H, s), 6.40 (1H, s), 7.42 (1H, m), 7.51 (2H, m), 7.76 (2H, d, J=7.2 Hz), 7.84 (2H, d, J=8.1 Hz), 7.97 (1H, s), 8.06 (2H, d, J=8.4 Hz), 8.65 (1H, s), 10.39 (1H, s).

20 FABMS(pos) 384.2 [M+H]⁺

Melting point: 202 - 203°C.

Example 134

25 4'-Chloro-N-[7-[(dimethylamino)methyl]-5,6-dihydro-3-quinolinyl][1,1'-biphenyl]-4-carboxamide



30 The titled compound was obtained by carrying out the same operation as in Example 1, using N-[(3-amino-5,6-dihydro-7-quinolinyl)methyl]-N,N-dimethylamine obtained

in Reference Example 86 and 4'-chloro[1,1'-biphenyl]-4-carboxylic acid.

¹H NMR (DMSO-d₆) δ : 2.17 (6H, s), 2.31 (2H, t, J=8.1 Hz), 2.85 (2H, t, J=8.1 Hz), 2.99 (2H, s), 6.41 (1H, s), 7.57 (2H, d, J=8.4 Hz), 7.81 (2H, d, J=8.4 Hz), 7.86 (2H, d, J=8.4 Hz), 7.98 (1H, s), 8.08 (2H, d, J=8.4 Hz), 8.66 (1H, s), 10.41 (1H, s).

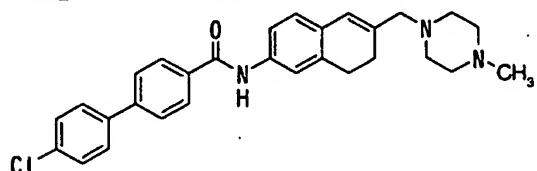
FABMS(pos) 418.2 [M+H]⁺

Melting point: 220 - 222°C.

10

Example 135

4'-Chloro-N-[6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



15 The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.

1H-NMR (CDCl₃) δ : 2.30 (3H, s), 2.25-2.50 (10H, m), 2.83 (2H, t, J = 8.1 Hz), 3.07 (2H, s), 6.35 (1H, s), 7.01 (1H, d, J = 8.1 Hz), 7.36 (1H, d, J = 7.8 Hz), 7.44 (2H, d, J = 8.4 Hz), 7.51 (1H, s), 7.55 (2H, d, J = 8.4 Hz), 7.66 (2H, d, J = 8.4 Hz), 7.84 (1H, s), 7.93 (2H, d, J = 8.4 Hz).

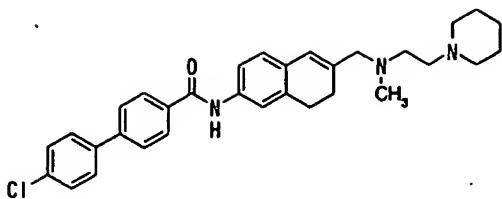
20 Melting point: 220 - 222°C (crystallization solvent : tetrahydrofuran - n-hexane)

25

Example 136

4'-Chloro-N-[6-[[methyl[2-(1-piperidinyl)ethyl]amino]methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

30



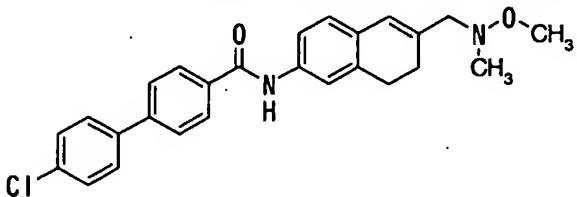
The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloromethyl-6-(chloromethyl)-7,8-dihydro-2-naphthalenyl[1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.

¹H-NMR (CDCl₃) δ : 1.72-1.77 (6H, m), 2.25-2.36 (2H, m), 2.27 (3H, s), 2.52-2.63 (8H, m), 2.84 (2H, t, J = 8.0 Hz), 3.08 (2H, s), 6.35 (1H, s), 7.01 (1H, d, J = 8.1 Hz), 7.38 (1H, d, J = 8.1 Hz), 7.44 (2H, d, J = 8.4 Hz), 7.49 (1H, s), 7.55 (2H, d, J = 8.4 Hz), 7.66 (2H, d, J = 8.4 Hz), 7.83 (1H, s), 7.93 (2H, d, J = 8.4 Hz).

Melting point: 165 - 167°C (crystallization solvent : tetrahydrofuran - n-hexane)

15 Example 137

4'-Chloro-N-[6-[[methoxy(methyl)amino]methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



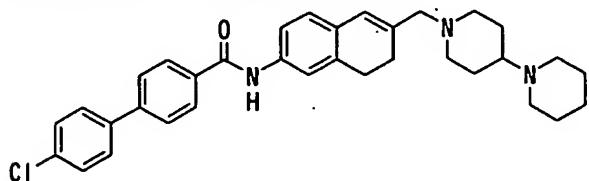
The titled compound was obtained by carrying out the same operation as in Example 51, using 4'-chloromethyl-6-(methoxy(methyl)amino)methyl-7,8-dihydro-2-naphthalenyl[1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.

¹H-NMR (CDCl₃) δ : 2.41 (2H, t, J = 8.1 Hz), 2.61 (3H, s), 2.86 (2H, t, J = 8.1 Hz), 3.37 (2H, s), 3.52 (3H, s), 6.39 (1H, s), 7.03 (1H, d, J = 8.1 Hz), 7.36 (1H, d, J = 8.1 Hz), 7.44 (2H, d, J = 8.4 Hz), 7.53 (1H, s), 7.55 (2H, d, J = 8.4 Hz), 7.66 (2H, d, J = 8.4 Hz), 7.83 (1H, s), 7.93 (2H, d, J = 8.4 Hz).

Melting point: 190 - 192°C (crystallization solvent : ethyl acetate - n-hexane)

Example 138

5 4'-Chloro-N-[6-[[4-(1-piperidinyl)-1-piperidinyl]methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

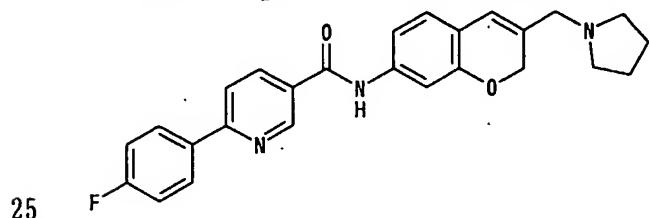


The titled compound was obtained by carrying out the
10 same operation as in Example 51, using 4'-chloro-N-[6-(chloromethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 56.
¹H-NMR (CDCl₃) δ : 1.45-1.96 (12H, m), 2.29-2.34 (3H, m), 2.57 (4H, s), 2.83 (2H, t, J = 8.1 Hz), 2.96-3.03 (4H, m),
15 6.32 (1H, s), 7.00 (1H, d, J = 8.1 Hz), 7.38 (1H, d, J = 8.1 Hz), 7.44 (2H, d, J = 8.4 Hz), 7.50 (1H, s), 7.55 (2H, d, J = 8.4 Hz), 7.66 (2H, d, J = 8.4 Hz), 7.86 (1H, s), 7.93 (2H, d, J = 8.4 Hz).

Melting point: 232 - 234°C (crystallization solvent :
20 ethyl acetate - n-hexane)

Example 139

6-(4-Fluorophenyl)-N-[3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]nicotineamide



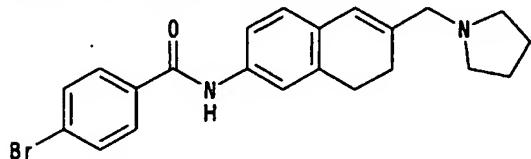
The titled compound was obtained by carrying out the
same operation as in Example 1, using 3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in
Reference Example 87.

¹H-NMR (CDCl₃) δ : 1.70 (4H, s), 2.43 (4H, s), 3.12 (2H, s), 4.73 (2H, s), 6.37 (1H, s), 7.03 (1H, d, J = 7.8 Hz), 7.29-7.40 (4H, m), 8.15 (1H, d, J = 8.4 Hz), 8.22-8.39 (3H, m), 9.15 (1H, s), 10.40 (1H, s).

5 Melting point: 233 - 235°C (crystallization solvent : tetrahydrofuran - n-hexane)

Example 140

10 4-Bromo-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]benzamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine

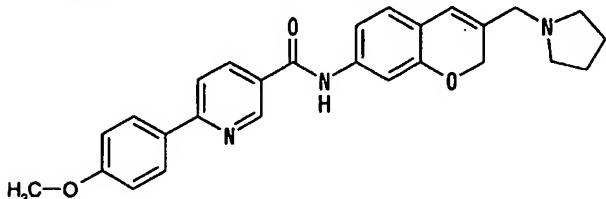
15 obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ : 1.79 (4H, s), 2.35 (2H, t, J = 8.1 Hz), 2.52 (4H, s), 2.83 (2H, t, J = 8.1 Hz), 3.17 (2H, s), 6.35 (1H, s), 6.99 (1H, d, J = 8.1 Hz), 7.34 (1H, d, J = 8.1 Hz), 7.43 (1H, s), 7.60 (2H, d, J = 8.4 Hz), 7.72 (2H, d, J = 8.4 Hz), 7.76 (1H, s).

20 Melting point: 135 - 137°C (crystallization solvent : ethyl acetate - n-hexane)

Example 141

25 6-(4-Methoxyphenyl)-N-[3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]nicotinamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 3-(1-

pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 87.

¹H-NMR (CDCl₃) δ : 1.70 (4H, s), 2.44 (4H, s), 3.12 (2H, s), 3.84 (3H, s), 4.73 (2H, s), 6.37 (1H, s), 7.03 (1H, d, J = 8.1 Hz), 7.09 (2H, t, J = 8.7 Hz), 7.29 (1H, d, J = 8.4 Hz), 7.31 (1H, s), 8.07 (1H, d, J = 8.7 Hz), 8.16 (2H, d, J = 8.7 Hz), 8.32 (1H, d, J = 8.4 Hz), 9.12 (1H, s), 10.34 (1H, s).

Elemental analysis for C₂₇H₂₇N₃O₃

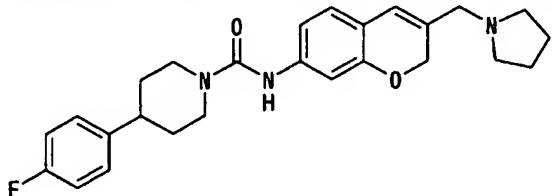
10 Calcd. : C, 73.45; H, 6.16; N, 9.52.

Found : C, 73.02; H, 6.27; N, 9.33.

Melting point: 243 - 245°C (crystallization solvent : tetrahydrofuran - n-hexane)

15 Example 142

4-(4-Fluorophenyl)-N-[3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]-1-piperidinecarboxamide



The titled compound was obtained by carrying out the same operation as in Example 99, using 3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 87.

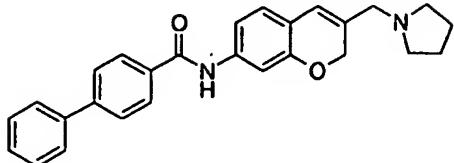
¹H-NMR (CDCl₃) δ : 1.69- 1.91 (8H, m), 2.49 (4H, s), 2.70 (1H, t, J = 12.0 Hz), 2.97 (2H, t, J = 12.0 Hz), 3.12 (2H, s), 4.19 (2H, d, J = 13.0 Hz), 4.76 (2H, s), 6.26 (1H, s), 6.37 (1H, s), 6.82-7.03 (5H, m), 7.16 (2H, dd, J = 5.4, 8.4 Hz).

Melting point: 176 - 178°C (crystallization solvent : ethyl acetate - diisopropyl ether)

30

Example 143

N-[3-(1-Pyrrolidinylmethyl)-2H-chromen-7-yl][1,1'-

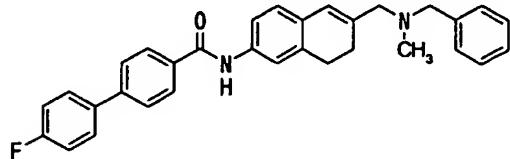
biphenyl]-4-carboxamide

The titled compound was obtained by carrying out the same operation as in Example 1, using 3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 87.

¹H-NMR (CDCl₃) δ : 1.79 (4H, s), 2.50 (4H, s), 3.15 (2H, s), 4.81 (2H, s), 6.30 (1H, s), 6.95 (1H, d, J = 8.1 Hz), 7.13 (1H, s), 7.20 (1H, d, J = 8.1 Hz), 7.39-7.50 (3H, m), 7.61-7.70 (4H, m), 7.82 (1H, s), 7.92 (2H, d, J = 8.1 Hz).
Melting point: 198 - 200°C (crystallization solvent : ethyl acetate)

Example 144

15 N-[6-[(N-Benzyl-N-methylamino)methyl]-7,8-dihydro-2-naphthalenyl]-4'-fluoro[1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N-benzyl-N-methylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 88.

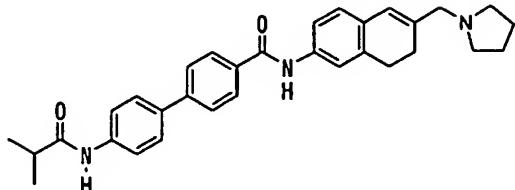
¹H-NMR (CDCl₃) δ : 2.20 (3H, s), 2.38 (2H, t, J = 8.1 Hz), 2.85 (2H, t, J = 8.1 Hz), 3.09 (2H, s), 3.52 (2H, s), 6.39 (1H, s), 7.02 (1H, d, J = 8.1 Hz), 7.13-7.66 (13H, m), 7.84 (1H, s), 7.93 (2H, d, J = 8.4 Hz).

Melting point: 143 - 145°C (crystallization solvent : ethyl acetate - n-hexane)

Example 145

30 4'-Isobutyrylamino-N-[6-(1-pyrrolidinylmethyl)-7,8-

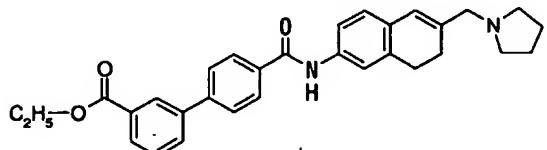
dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as an amorphous powder by carrying out the same operation as in Example 1, 5 using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54. MS m/z 494.4 (MH⁺).

Example 146

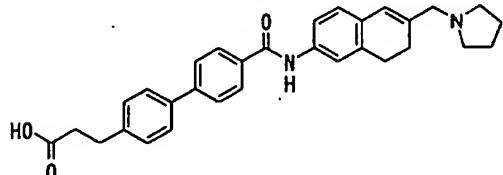
10 Ethyl 4'-[[[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]amino]carbonyl][1,1'-biphenyl]-3-carboxylate



The titled compound was obtained as an amorphous powder by carrying out the same operation as in Example 1, 15 using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54. MS m/z 481.4 (MH⁺).

20 Example 147

3-[4'-[[[6-(1-Pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]amino]carbonyl][1,1'-biphenyl]-4-yl]propionic acid

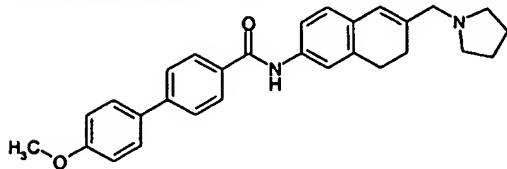


25 The titled compound was obtained as a powder by carrying out the same operation as in Example 1, using

6-(1-pyrrolidinylmethyl)-7,8--dihydro-2-naphthalenamine obtained in Reference Example 54.
MS m/z 481.4 (MH⁺).

5 Example 148

4'-Methoxy-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the
10 same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ : 1.80 (4H, m), 2.36 (2H, t, J=7.8 Hz), 2.52 (4H, m), 2.86 (2H, t, J=7.8 Hz), 3.18 (2H, s), 3.87 (3H, s), 6.36 (1H, s), 7.00-7.03 (3H, m), 7.26 (1H, m), 7.38 (1H, d, J=8.3 Hz), 7.49 (1H, s), 7.58 (2H, d, J=8.6 Hz), 7.67 (1H, d, J=8.2 Hz), 7.78 (1H, s), 7.90 (2H, d, J=8.2 Hz).

Elemental analysis for C₂₉H₃₀N₂O₂

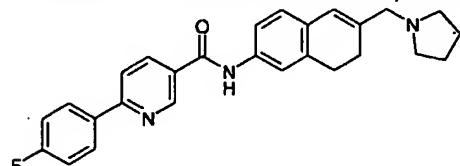
Calcd. : C, 79.42; H, 6.89; N, 6.39.

20 Found : C, 79.21; H, 6.88; N, 6.35.

Melting point: 187-188 °C (crystallization solvent : ethyl acetate - diisopropyl ether)

Example 149

25 6-(4-Fluorophenyl)-N-[6-[(1-pyrrolidinyl)methyl]-7,8-dihydro-2-naphthalenyl]nicotinamide



The titled compound was obtained by carrying out the
same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine

obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ : 1.81 (4H, m), 2.36 (2H, t, J=8.1 Hz), 2.53 (4H, m), 2.86 (2H, t, J=8.1 Hz), 3.18 (2H, s), 6.37 (1H, s), 7.03 (1H, d, J=7.8 Hz), 7.16-7.30 (3H, m), 7.47 (1H, s), 7.77-7.82 (2H, m), 8.06 (2H, dd, J=8.9, 5.3 Hz), 8.25 (1H, dd, J=8.4, 2.2 Hz), 9.11 (1H, d, J=2.0 Hz).

Elemental analysis for C₂₇H₂₆FN₃O

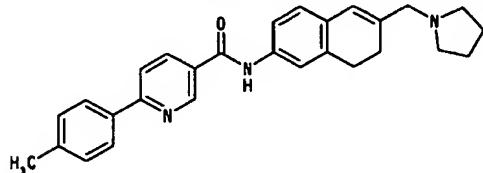
Calcd.: C, 75.85; H, 6.13; N, 9.83.

Found: C, 75.71; H, 5.93; N, 9.75.

10 Melting point: 225-227 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 150

15 6-(4-Methylphenyl)-N-[6-[(1-pyrrolidinyl)methyl]-7,8-dihydro-2-naphthalenyl]nicotinamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine

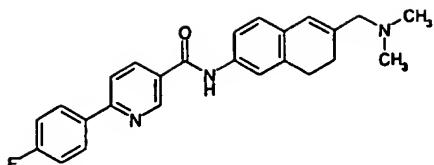
20 obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ : 1.81 (4H, m), 2.36 (2H, t, J=7.8 Hz), 2.43 (3H, s), 2.53 (4H, m), 2.86 (2H, t, J=7.8 Hz), 3.19 (2H, s), 6.37 (1H, s), 7.02 (1H, d, J=8.7 Hz), 7.25-7.39 (3H, m), 7.47 (1H, s), 7.82 (2H, m), 7.96 (2H, d, J=8.1 Hz), 8.23 (1H, dd, J=8.1, 2.3 Hz), 9.12 (1H, d, J=2.3 Hz).

25 Melting point: 235-236 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 151

30 N-[6-[(Dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]-6-(4-fluorophenoxy)nicotinamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 41-2).

¹H-NMR (CDCl₃) δ : 2.25 (6H, s), 2.34 (2H, t, J=8.1 Hz), 2.86 (2H, t, J=8.1 Hz), 2.99 (2H, s), 6.35 (1H, s), 7.03 (1H, d, J=8.1 Hz), 7.17 (2H, m), 7.26 (1H, m), 7.39 (1H, d, J=8.1 Hz), 7.47 (1H, s), 7.78 (1H, d, J=7.2 Hz), 7.83 (1H, s), 8.06 (1H, dd, J=8.4, 6.7 Hz), 8.25 (1H, d, J=6.7 Hz), 9.12 (1H, s).

Elemental analysis for C₂₅H₂₄FN₃O

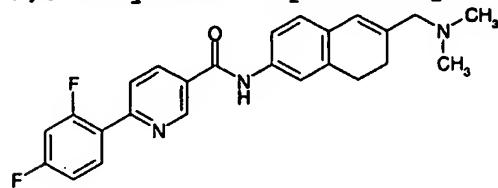
Calcd. : C, 74.79; H, 6.03; N, 10.47.

Found : C, 74.74; H, 5.95; N, 10.24.

Melting point: 216-219 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 152

6-(2,4-Difluorophenyl)-N-[6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]nicotinamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 41-2).

¹H-NMR (CDCl₃) δ : 2.25 (6H, s), 2.34 (2H, t, J=8.1 Hz), 2.85 (2H, t, J=8.1 Hz), 3.00 (2H, s), 6.35 (1H, s), 6.90-7.06 (3H, m), 7.39 (1H, d, J=7.8 Hz), 7.47 (1H, s), 7.80-7.90 (2H, m), 8.10 (1H, dd, J=15.3, 8.8 Hz), 8.23 (1H, dd, J=8.4, 2.3 Hz), 9.15 (1H, d, J=1.7 Hz).

Elemental analysis for C₂₅H₂₃F₂N₃O

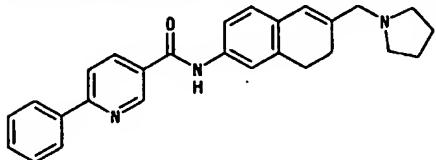
Calcd. : C, 71.58; H, 5.53; N, 10.02.

Found : C, 71.50; H, 5.49; N, 9.61.

Melting point: 162-163 °C (crystallization solvent: ethyl
5 acetate - diisopropyl ether)

Example 153

6-Phenyl-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide



10

The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

15

¹H-NMR (CDCl₃) δ : 1.81 (4H, m), 2.36 (2H, t, J=8.1 Hz), 2.53 (4H, m), 2.85 (2H, t, J=8.1 Hz), 3.18 (2H, s), 6.37 (1H, s), 7.02 (1H, d, J=8.1 Hz), 7.37-7.53 (5H, m), 7.83 (1H, d, J=8.1 Hz), 7.86 (1H, d, J=6.2 Hz), 8.04 (1H, s), 8.06 (1H, d, J=1.7 Hz), 8.24 (1H, dd, J=8.4, 2.4 Hz), 9.13 (1H, d, J=2.2 Hz).

20

Elemental analysis for C₂₇H₂₇N₃O

Calcd. : C, 79.19; H, 6.65; N, 10.26.

Found : C, 78.93; H, 6.65; N, 10.19.

25

Melting point: 186-187 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 154

6-(4-Methoxyphenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide

30



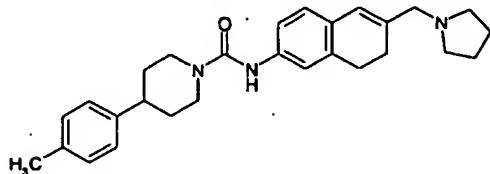
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

5 ¹H-NMR (CDCl₃) δ : 1.80 (4H, m), 2.36 (2H, t, J=8.1 Hz), 2.52 (4H, m), 2.84 (2H, t, J=8.1 Hz), 3.18 (2H, s), 3.88 (3H, s), 6.36 (1H, s), 7.02 (3H, m), 7.37 (1H, d, J=7.5 Hz), 7.47 (1H, s), 7.78 (1H, d, J=8.1 Hz), 7.79 (1H, s), 8.03 (2H, d, J=8.5 Hz), 8.20 (1H, d, J=8.1 Hz), 9.08 (1H, s).

10 Melting point: 219-220 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 155

4-(4-Methylphenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

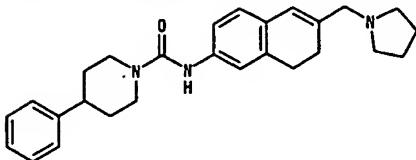
20 ¹H-NMR (CDCl₃) δ : 1.64-1.92 (8H, m), 2.29 (2H, m), 2.32 (3H, s), 2.51 (4H, m), 2.64 (1H, m), 2.80 (2H, t, J=7.8 Hz), 2.97 (2H, dd, J=13.1, 10.7 Hz), 3.15 (2H, s), 4.19 (2H, d, J=13.1 Hz), 6.32 (1H, s), 6.35 (1H, s), 6.42 (1H, d, J=7.8 Hz), 7.06-7.20 (6H, m)

Elemental analysis for C₂₈H₃₅N₃O · 0.5H₂O
Calcd.: C, 76.67; H, 8.27; N, 9.58.
Found: C, 76.72; H, 8.03; N, 9.36.
Melting point: 197-198 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 156

4-Phenyl-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-

naphthalenyl]-1-piperidinecarboxamide



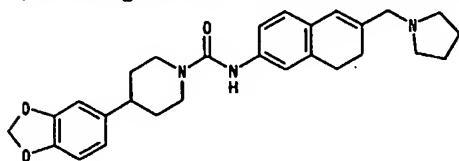
The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ : 1.72-1.94 (8H, m), 2.32 (2H, t, J=8.1 Hz), 2.50 (4H, m), 2.72 (1H, m), 2.80 (2H, t, J=8.1 Hz), 2.99 (2H, dd, J=13.4, 10.6 Hz), 3.16 (2H, s), 4.21 (2H, d, J=13.4 Hz), 6.32 (1H, s), 6.34 (1H, s), 6.93 (1H, d, J=8.4 Hz), 7.07 (1H, d, J=8.1 Hz), 7.20-7.35 (6H, m).

Melting point: 184-186 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

15 Example 157

4-(1,3-Benzodioxol-5-yl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



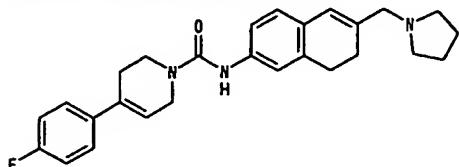
The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ : 1.61-1.88 (8H, m), 2.31 (2H, t, J=8.1 Hz), 2.51 (4H, m), 2.59 (1H, m), 2.62 (2H, t, J=8.1 Hz), 2.94 (2H, dd, J=13.1, 11.2 Hz), 3.15 (2H, s), 4.18 (2H, d, J=13.1 Hz), 5.93 (2H, s), 6.31 (1H, s), 6.44 (1H, s), 6.64-6.77 (3H, m), 6.92 (1H, d, J=8.1 Hz), 7.07 (1H, d, J=8.1 Hz), 7.19 (1H, s).

Melting point: 149-150 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 158

4-(4-Fluorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-3,6-dihydro-1(2H)-
5 pyridinecarboxamide



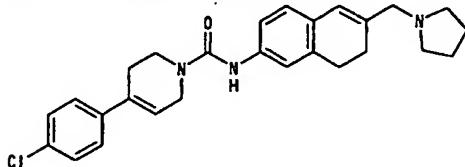
The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine
10 obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ : 1.79 (4H, m), 2.32 (2H, t, J=8.1 Hz), 2.50 (4H, m), 2.59 (2H, brt), 2.80 (2H, t, J=8.1 Hz), 3.17 (2H, s), 3.74 (2H, t, J=5.7 Hz), 4.15 (2H, d, J=2.5 Hz), 6.00 (1H, brt), 6.32 (1H, s), 6.32 (1H, s), 6.94 (1H, d, J=8.1 Hz), 7.00-7.32 (6H, m).

Elemental analysis for C₂₇H₃₀FN₃O
Calcd. : C, 75.15; H, 7.01; N, 9.74.
Found : C, 75.09; H, 6.93; N, 9.77.
Melting point: 206-207 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 159

4-(4-Chlorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-3,6-dihydro-1(2H)-
25 pyridinecarboxamide



The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine
30 obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ : 1.79 (4H, m), 2.32 (2H, t, J=8.1 Hz), 2.50 (4H, m), 2.59 (2H, brt), 2.80 (2H, t, J=8.1 Hz), 3.16 (2H, s), 3.73 (2H, t, J=5.6 Hz), 4.15 (2H, d, J=2.8 Hz), 6.06 (1H, brt), 6.30 (1H, s), 6.32 (1H, s), 6.93 (1H, d, J=7.8 Hz), 7.09 (1H, d, J=7.8 Hz), 7.21-7.31 (5H, m).

Elemental analysis for C₂₇H₃₀ClN₃O

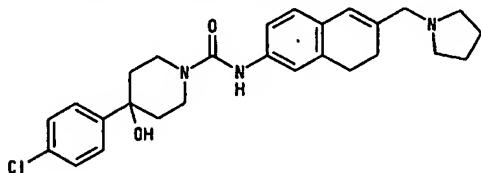
Calcd. : C, 72.39; H, 6.75; N, 9.38.

Found : C, 72.19; H, 6.75; N, 9.19.

Melting point: 217-218 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 160

4-(4-Chlorophenyl)-4-hydroxy-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



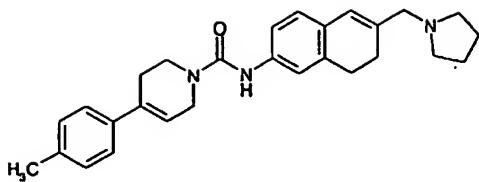
The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ : 1.79 (4H, m), 1.80 (2H, m), 2.04 (1H, dd, J=13.1, 10.8 Hz), 2.06 (1H, dd, J=13.1, 10.8 Hz), 2.31 (2H, t, J=7.8 Hz), 2.50 (1H, brs), 2.51 (4H, m), 2.79 (2H, t, J=7.8 Hz), 3.15 (2H, s), 3.41 (2H, dd, J=12.6, 10.8 Hz), 4.00 (2H, d, J=12.6 Hz), 6.32 (1H, s), 6.37 (1H, s), 6.93 (1H, d, J=8.1 Hz), 7.05-7.42 (6H, m).

Melting point: 181-182 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 161

4-(4-Methylphenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-3,6-dihydro-1(2H)-pyridinecarboxamide



The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine

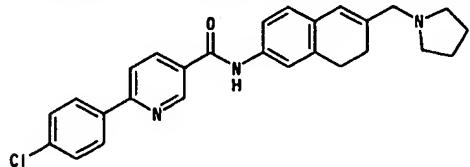
5 obtained in Reference Example 54.

¹H-NMR (CDCl₃) δ : 1.79 (4H, m), 2.32 (2H, t, J=7.8 Hz), 2.35 (3H, s), 2.50 (4H, m), 2.61 (2H, brt), 2.80 (2H, t, J=7.8 Hz), 3.16 (2H, s), 3.73 (2H, t, J=5.7 Hz), 4.15 (2H, d, J=2.8 Hz), 6.03 (1H, s), 6.29 (1H, s), 6.32 (1H, s), 6.93 (1H, d, J=8.1 Hz), 7.07-7.30 (6H, m).

10 Melting point: 199-202 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 162

15 6-(4-Chlorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine

20 obtained in Reference Example 54.

¹H-NMR (CDCl₃+DMSO-d₆) δ : 1.80 (4H, m), 2.32-2.58 (6H, m), 2.85 (2H, t, J=8.0 Hz), 3.18 (2H, s), 6.36 (1H, s), 7.01 (1H, d, J=8.4 Hz), 7.48 (2H, d, J=8.4 Hz), 7.49 (1H, m), 25 7.59 (1H, s), 7.83 (1H, d, J=8.4 Hz), 8.04 (2H, d, J=8.4 Hz), 8.35 (1H, dd, J=8.4, 2.2 Hz), 9.25 (1H, d, J=2.2 Hz), 9.42 (1H, s).

Elemental analysis for C₂₂H₂₆ClN₃O

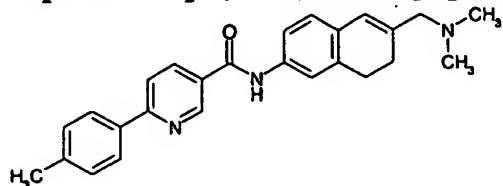
Calcd.: C, 73.04; H, 5.90; N, 9.46.

30 Found: C, 73.11; H, 5.71; N, 9.20.

Melting point: 252-253 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 163

5 N-[6-[(Dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]-6-(4-methylphenyl)nicotinamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 41-2).

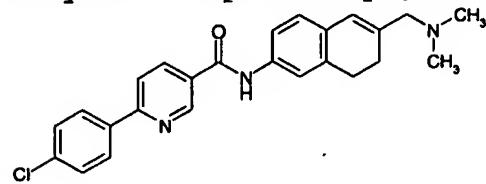
10 ¹H-NMR (CDCl₃) δ: 2.25 (6H, s), 2.34 (2H, t, J=8.1 Hz), 2.43 (3H, s), 2.85 (2H, t, J=8.1 Hz), 2.99 (2H, s), 6.34 (1H, s), 7.02 (1H, d, J=8.1 Hz), 7.31 (2H, d, J=8.1 Hz), 15 7.39 (1H, d, J=8.1 Hz), 7.46 (1H, s), 7.81 (1H, d, J=8.4 Hz), 7.87 (1H, s), 7.96 (2H, d, J=8.1 Hz), 8.22 (1H, dd, J=8.4, 2.3 Hz), 9.11 (1H, d, J=2.3 Hz).

Melting point: 228-230 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

20

Example 164

6-(4-Chlorophenyl)-N-[6-[(dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]nicotinamide



25 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 41-2).

30 ¹H-NMR (CDCl₃) δ: 2.25 (6H, s), 2.35 (2H, t, J=8.1 Hz), 2.86 (2H, t, J=8.1 Hz), 2.99 (2H, s), 6.35 (1H, s), 7.04 (1H,

d, J=8.1 Hz), 7.40 (1H, d, J=8.4 Hz), 7.49 (1H, brs), 7.49 (2H, d, J=8.4 Hz), 7.78 (1H, s), 7.84 (1H, d, J=8.4 Hz), 8.02 (2H, d, J=8.4 Hz), 8.26 (1H, dd, J=8.1, 2.2 Hz), 9.13 (1H, d, J=2.2 Hz).

5 Elemental analysis for C₂₅H₂₄ClN₃O

Calcd. : C, 71.85; H, 5.79; N, 10.05.

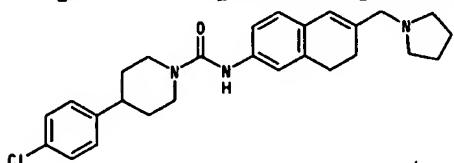
Found: C, 71.88; H, 5.67; N, 9.86.

Melting point: 248-249 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

10

Example 165

4-(4-Chlorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



15 The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

1H-NMR (CDCl₃) δ : 1.66-1.91 (8H, m), 2.32 (2H, t, J=8.1 Hz), 2.50 (4H, m), 2.70 (1H, m), 2.80 (2H, t, J=8.1 Hz), 2.98 (2H, dd, J=13.7, 12.0 Hz), 3.16 (2H, s), 4.20 (2H, d, J=13.7 Hz), 6.32 (1H, s), 6.32 (1H, s), 6.93 (1H, d, J=8.1 Hz), 7.05-7.30 (6H, m).

Elemental analysis for C₂₇H₃₂ClN₃O

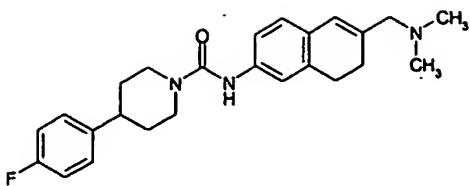
Calcd. : C, 72.06; H, 7.17; N, 9.34.

Found: C, 72.08; H, 7.23; N, 9.15.

Melting point: 194-195 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

30 Example 166

N-[6-[(Dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]-4-(4-fluorophenyl)-1-piperidinecarboxamide

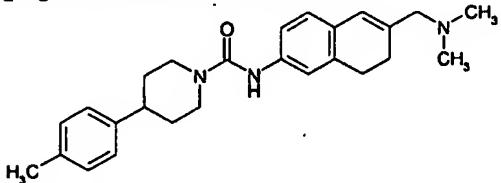


The titled compound was obtained by carrying out the same operation as in Example 99, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 41-2).

¹H-NMR (CDCl₃) δ: 1.65-1.75 (2H, m), 1.89 (2H, d, J=11.4 Hz), 2.23 (6H, s), 2.30 (2H, t, J=8.1 Hz), 2.70 (1H, m), 2.80 (2H, t, J=8.1 Hz), 2.94-3.01 (4H, m), 4.20 (2H, d, J=13.4 Hz), 6.30 (1H, s), 6.35 (1H, s), 6.92-7.20 (7H, m).
10 Melting point: 187-188 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 167

N-[6-[(Dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl]-4-(4-methylphenyl)-1-piperidinecarboxamide
15



The titled compound was obtained by carrying out the same operation as in Example 99, using 6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 41-2).

¹H-NMR (CDCl₃) δ: 1.66-1.74 (2H, m), 1.89 (2H, d, J=11.7 Hz), 2.28 (6H, s), 2.30 (2H, t, J=8.1 Hz), 2.38 (3H, s), 2.68 (1H, m), 2.80 (2H, t, J=8.1 Hz), 2.94-3.02 (4H, m), 25 4.19 (2H, d, J=12.8 Hz), 6.30 (1H, s), 6.35 (1H, s), 6.93 (1H, d, J=8.1 Hz), 7.07-7.20 (6H, m).

Elemental analysis for C₂₆H₃₃N₃O · 0.5H₂O

Calcd.: C, 75.69; H, 8.31; N, 10.18

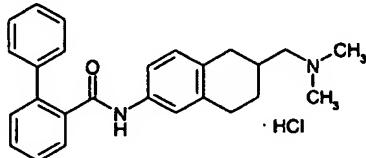
Found: C, 75.44; H, 8.16; N, 10.05

30 Melting point: 200-202 °C (crystallization solvent: ethyl

acetate - diisopropyl ether)

Example 168

5 N-[6-[(Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-2-carboxamide hydrochloride



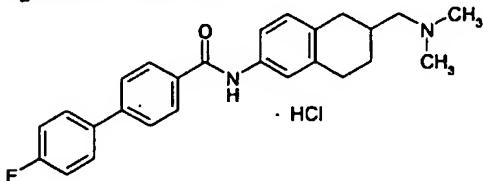
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-amino-2-[(N,N-dimethylamino)methyl]tetralin hydrochloride.

¹H-NMR (DMSO-d₆) δ: 1.39 (1H, m), 1.99 (1H, m), 2.17 (1H, m), 2.42 (1H, dd, J=16.2, 10.1 Hz), 2.78 (6H, s), 2.88 (1H, dd, J=16.2, 4.5 Hz), 3.06 (2H, t, J=5.7 Hz), 3.38 (2H, s), 6.94-7.62 (11H, m), 7.64 (1H, d, J=1.7 Hz), 10.11 (1H, brs), 15 10.18 (1H, s).

Melting point: 196-197 °C (crystallization solvent: methanol - ethyl acetate)

Example 169

20 N-[6-[(Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-fluoro[1,1'-biphenyl]-4-carboxamide hydrochloride



25 4'-Fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide synthesized in Example 42 was dissolved in ethyl acetate. An excess amount of 4N hydrochloric acid-ethyl acetate solution was added to the solution, which was concentrated under reduced pressure. The 30 resulting residue was recrystallized from methanol - ethyl

acetate, to give the titled compound.

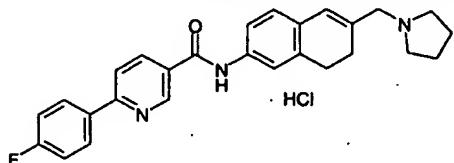
¹H-NMR (DMSO-d₆) δ : 1.43 (1H, m), 2.06 (1H, m), 2.21 (1H, m), 2.45 (1H, m), 2.79 (6H, s), 2.92 (1H, dd, J=16.2, 4.2 Hz), 3.08 (2H, d, J=6.4 Hz), 3.33 (2H, s), 7.05 (1H, d, J=8.4 Hz), 7.34 (2H, dd, J=8.9, 8.9 Hz), 7.53 (1H, d, J=8.4 Hz), 7.59 (1H, s), 7.80 (4H, m), 8.06 (2H, d, J=8.1 Hz), 10.02 (1H, s), 10.03 (1H, brs).

Melting point: 240-245 °C (crystallization solvent: methanol - ethyl acetate)

10

Example 170

6-(4-Fluorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide hydrochloride



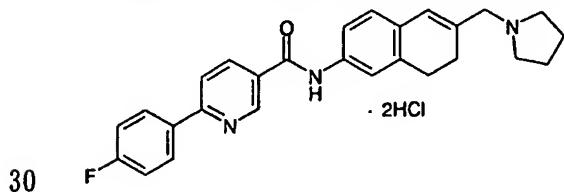
15 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

¹H-NMR (DMSO-d₆) δ : 1.70 (4H, m), 2.26 (2H, t, J=8.1 Hz), 2.44 (4H, m), 2.76 (2H, t, J=8.1 Hz), 3.12 (2H, s), 3.34 (1H, s), 6.36 (1H, s), 7.03 (1H, d, J=7.8 Hz), 7.37 (2H, dd, J=8.4, 7.0 Hz), 7.57 (1H, d, J=8.4 Hz), 7.59 (1H, s), 8.13-8.42 (4H, m), 9.19 (1H, s), 10.43 (1H, s).

Melting point: 229-231 °C (crystallization solvent: methanol - ethyl acetate)

Example 171

6-(4-Fluorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide dihydrochloride



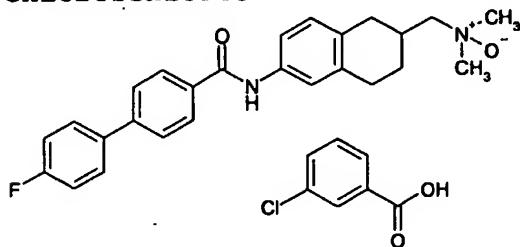
The titled compound was obtained by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

5 ¹H-NMR (DMSO-d₆) δ : 2.00 (4H, m), 2.45 (4H, m), 2.83 (2H, t, J=8.1 Hz), 3.05 (2H, m), 3.47 (2H, m), 3.88 (1H, s), 6.69 (1H, s), 7.13 (1H, d, J=8.1 Hz), 7.38 (2H, dd, J=8.9, 8.6 Hz), 7.64 (1H, d, J=10.6 Hz), 7.66 (1H, s), 8.14-8.42 (4H, m), 9.19 (1H, s), 10.52 (1H, s), 10.60 (1H, brs).

10 Melting point: 245-248 °C (crystallization solvent: methanol - ethyl acetate)

Example 172

15 N-[6-[(Dimethylnitrooyl)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-4'-fluoro[1,1'-biphenyl]-4-carboxamide 3-chlorobenzoate

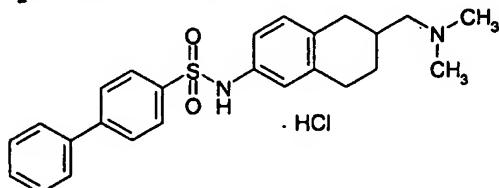


20 4'-FluoroN-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide (100 mg) obtained in Example 42 was dissolved in acetone (10 ml), which was stirred under ice-cooling. 3-Chloroperbenzoic acid (purity : 50%) (86 mg) was added to the solution, which was stirred under ice-cooling for 1 hour. The reaction mixture was concentrated under reduced pressure, and the residue was washed with diisopropyl ether, to give the titled compound (158 mg).
¹H-NMR (DMSO-d₆) δ : 1.57 (1H, m), 2.07 (1H, m), 2.61 (1H, m), 2.82 (2H, m), 3.04 (1H, m), 3.33 (1H, m), 3.48 (6H, s), 3.56-3.67 (2H, m), 6.55 (1H, s), 7.03 (1H, d, J=8.4 Hz), 7.30-7.56 (6H, m), 7.78-7.85 (6H, m), 8.04 (2H, d, J=8.4 Hz), 10.17 (1H, s).

FABMS(pos) 419.1 [M+H]⁺

Example 173

5 N-[6-[(Dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-sulfonamide hydrochloride



10 6-[(N, N-Dimethylamino)methyl]-7,8-dihydro-2-naphthalenamine (200 mg, 0.72 mmol) obtained in Example 41-2 was dissolved in acetonitrile (30 ml). Triethylamine (0.401 ml, 2.88 mmol) and [1,1'-biphenyl]-4-sulfonylchloride (200 mg, 0.79 mmol) were added to the solution under ice-cooling, which was stirred for 3 hours. The reaction mixture was concentrated. Ethyl 15 acetate and water were added to the residue, and extraction was conducted. The ethyl acetate layer was concentrated, and the residue was purified by alumina column chromatography (development solvent; ethyl acetate:n-hexane = 33:67). 4N Hydrogen chloride-ethyl acetate 20 solution was added to the resulting oily substance, which was concentrated. The residue was recrystallized from methanol - ethyl acetate, to give the titled compound (194 mg).

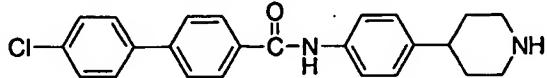
25 ¹H-NMR (DMSO-d₆) δ: 1.32 (1H, m), 1.96 (1H, m), 2.11 (1H, m), 2.35 (1H, d, J=15.9, 10.0 Hz), 2.74 (2H, m), 2.78 (7H, m), 3.02 (2H, m), 6.89 (2H, d, J=10.6 Hz), 6.91 (1H, m), 7.40-7.51 (3H, m), 7.70 (2H, d, J=6.7 Hz), 7.85 (4H, m), 9.92 (1H, brs), 10.23 (1H, s).

30 Melting point: 168-170 °C (crystallization solvent: methanol - ethyl acetate)

FABMS(pos) 421.1 [M+H]⁺

Example 174

4'-Chloro-N-[4-(4-piperidinyl)phenyl][1,1'-biphenyl]-4-carboxamide



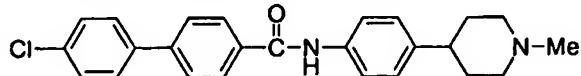
5 The titled compound was obtained as a colorless powder by carrying out the same operation as in Example 127-2), using 4'-chloro-N-[4-(4-piperidinyl)phenyl][1,1'-biphenyl]-4-carboxamide obtained in Reference Example 89.
¹H-NMR (CDCl₃+ DMSO-d₆) δ: 1.40-1.90 (4H, m), 2.60-2.90
10 (3H, m), 3.18-3.28 (2H, m), 7.19 (2H, d, J=8.1 Hz), 7.49 (2H, d, J=7.0 Hz), 7.67-7.75 (6H, m), 8.07-8.10 (3H, m), 10.16 (1H, s).

Melting point: 276-281 °C (decomposition)
crystallization solvent: ethyl acetate)

15

Example 175

4'-Chloro-N-[4-(1-methyl-4-piperidinyl)phenyl][1,1'-biphenyl]-4-carboxamide



20 A mixture of 4'-chloro-N-[4-(4-piperidinyl)phenyl][1,1'-biphenyl]-4-carboxamide (0.17 g) obtained in Example 174, 37% aqueous formaldehyde solution (0.05 ml) and formic acid (0.5 ml) was heated at 100°C for 4 hours. The reaction mixture was cooled to room
25 temperature. Water was added to the mixture, which was made alkaline with 8N aqueous sodium hydroxide solution, and extracted with ethyl acetate - tetrahydrofuran (1:1) mixed solution. The extract was washed with saturated aqueous sodium chloride solution, dried over anhydrous magnesium sulfate, and then the solvent was distilled out under reduced pressure. The resulting solid was washed with ethyl acetate, dried under reduced pressure, to give the
30 titled compound (90 mg).

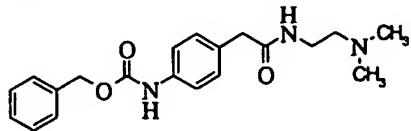
¹H-NMR (CDCl₃+ DMSO-d₆) δ: 1.55-1.80 (2H, m), 1.90-2.10

(2H, m), 2.22 (3H, s), 2.30-2.45 (1H, m), 2.80-3.20 (4H, m), 7.11 (2H, d, J=8.1 Hz), 7.36 (2H, d, J=8.1 Hz), 7.50-7.63 (6H, m), 7.97 (2H, d, J=8.4 Hz), 9.79 (1H, s).

Melting point: 273-277 °C (decomposition) (Washing solvent: ethyl acetate)

Example 176

Benzyl 4-[2-[(2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenylcarbamate



10

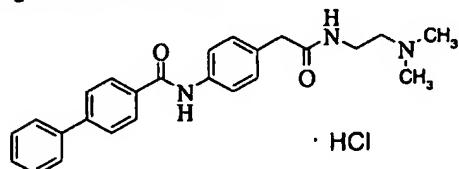
N,N-Dimethylethylenediamine (0.64 ml), WSC (1.31 g), HOBt (1.05 g), and triethylamine (2.4 ml) were added to a tetrahydrofuran (50 ml) solution of 2-[4-[(benzyloxy)carbonyl]amino]phenyl]acetic acid (1.5 g) obtained in Reference Example 90.. After stirring for 20 hours, the reaction mixture was poured into water, and extraction was conducted using ethyl acetate. The organic layer was washed with water, saturated aqueous sodium bicarbonate solution, and saturated aqueous sodium chloride solution, dried and then concentrated. The residue was recrystallized from ethyl acetate - hexane, to give the titled compound (1.72 g).

Melting point: 126-127 °C.

25

Example 177

N-[4-[2-[(2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



30

Oxalyl chloride (0.56 ml) was added dropwise to a

tetrahydrofuran (45 ml) solution of 4-biphenylcarboxylic acid (1.01 g) under ice-cooling. 9 drops of DMF was added to the mixture, and the temperature of the mixture was raised to room temperature, which was stirred for 40 minutes.

5 The reaction mixture was concentrated and dried.

A tetrahydrofuran (50 ml) solution of the residue was added dropwise to a tetrahydrofuran (45 ml) solution of 2-(4-aminophenyl)-N-[2-(dimethylamino)ethyl]acetamide (939 mg) obtained in Reference Example 91 under ice-cooling.

10 Then the temperature of the reaction mixture was raised to room temperature, which was stirred for 2 hours.

Saturated aqueous sodium bicarbonate solution was added to the reaction mixture, and extraction was conducted using ethyl acetate. The organic layer was washed with water and saturated aqueous sodium chloride solution, dried over sodium sulfate, and then concentrated. The residue was dissolved in tetrahydrofuran. 4N Hydrochloric acid-ethyl acetate was added to the solution, which was concentrated.

15 The residue was recrystallized from methanol - diisopropyl ether, to give the titled compound (750 mg). Melting point: 216-217 °C.

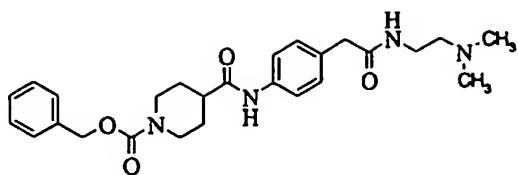
The above N-[4-[2-[(2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl][1,1'-biphenyl]-4-carboxyamide hydrochloride (100 mg) was dissolved in saturated aqueous sodium bicarbonate solution, and extraction was conducted using tetrahydrofuran-ethyl acetate (1:1). The organic layer was washed with saturated aqueous sodium chloride solution, dried over sodium sulfate, and then concentrated.

20 The residue was recrystallized from methanol - diisopropyl ether, to give a free base form (56 mg) of the titled compound.

Melting point: 228-229 °C.

Example 178

35 Benzyl 4-[[4-[2-[(2-(dimethylamino)ethyl]amino]-2-oxoethyl]anilino]carbonyl]-1-piperidinecarboxylate

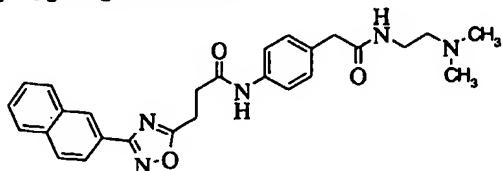


2-(4-Aminophenyl)-N-[2-(dimethylamino)ethyl]acetamide (221 mg), WSC (249 mg), 1-hydroxybenzotriazole (199 mg), triethylamine (0.4 ml),
 5 and dimethylaminopyridine (244 mg) were added to a tetrahydrofuran (10 ml) solution of 1-[(benzyloxy)carbonyl]-4-piperidinecarboxylic acid (290 mg), which was stirred for 20 hours. The reaction mixture was poured into water, and extraction was conducted using
 10 ethyl acetate. The organic layer was washed with water, saturated aqueous sodium bicarbonate solution, and saturated aqueous sodium chloride solution, dried over sodium sulfate, and then concentrated. The residue was recrystallized from methanol - diisopropyl ether, to give
 15 the titled compound (230 mg).

Melting point: 169-170 °C.

Example 179

N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-3-[3-(2-naphthyl)-1,2,4-oxadiazol-5-yl]propanamide



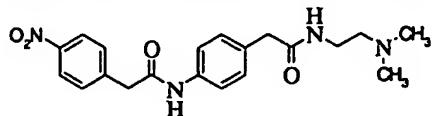
2-(4-Aminophenyl)-N-[2-(dimethylamino)ethyl]acetamide (221 mg), WSC (249 mg),
 25 1-hydroxybenzotriazole (199 mg), triethylamine (0.4 ml), and dimethylaminopyridine (244 mg) were added to a DMF (5 ml) solution of 3-[3-(2-naphthyl)-1,2,4-oxadiazol-5-yl]propionic acid (268 mg), which was stirred for 5 hours. The reaction mixture was poured into water, and extraction was conducted using ethyl acetate. The organic layer was

washed with water, saturated aqueous sodium bicarbonate solution, and saturated aqueous sodium chloride solution, dried over sodium sulfate, and then concentrated. The residue was recrystallized from methanol, to give the
5 titled compound (166 mg).

Melting point: 173-174 °C.

Example 180

10 N-[4-[2-[(2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-2-(4-nitrophenyl)acetamide



15 2-(4-Aminophenyl)-N-[2-(dimethylamino)ethyl]acetamide (221 mg), WSC (free form : 0.23 ml), 1-hydroxybenzotriazole (199 mg), and dimethylaminopyridine (244 mg) were added to a DMF (5 ml) solution of 4-nitrophenylacetic acid (181 mg), which was stirred for 4 hours. The reaction mixture was poured into water, and extraction was conducted using ethyl acetate. The organic layer was washed with water, saturated aqueous sodium bicarbonate solution, and saturated aqueous sodium chloride solution, dried over sodium sulfate, and then concentrated. The residue was recrystallized from methanol, to give the titled compound (80 mg).

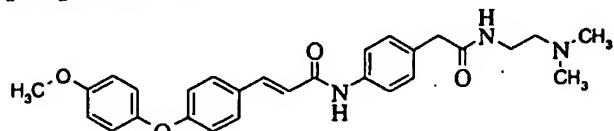
Melting point: 160-162 °C.

25

Example 181

(E)-N-[4-[2-[(2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-3-[4-(4-methoxyphenoxy)phenyl]-2-propanamide

30



2-(4-Aminophenyl)-N-[2-(dimethylamino)ethyl]acetamide (221 mg), WSC (free form :

0.23 ml), 1-hydroxybenzotriazole (199 mg), triethylamine (0.14 ml) and dimethylaminopyridine (122 mg) were added to a DMF (5 ml) solution of (E)-3-[4-(4-methoxyphenoxy)phenyl]-2-propenoic acid (270 mg), which
 5 was stirred for 24 hours. The reaction mixture was poured into water, and extraction was conducted using ethyl acetate - tetrahydrofuran (1:1). The organic layer was washed with water, saturated aqueous sodium bicarbonate solution, and saturated aqueous sodium chloride solution,
 10 dried over sodium sulfate, and then concentrated. The resulting crude crystals were washed with diisopropyl ether, to give the titled compound (227 mg).

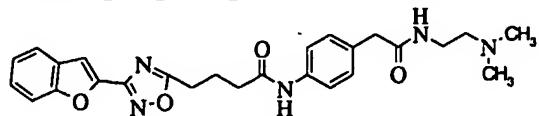
Melting point: 175-177 °C (decomposition).

15 Compounds described in the following Example 182 to 198 were produced in the same manner as in Example 181.

Example 182

4-[3-(1-Benzofuran-2-yl)-1,2,4-oxadiazol-5-yl]-N-[4-[2-[(2-(dimethylamino)ethyl]amino]-2-

20 oxoethyl]phenyl]butanamide

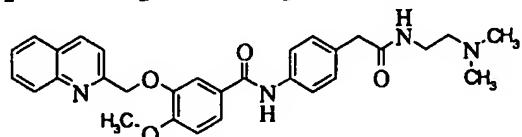


Melting point: 161-163 °C.

Washing solvent: diisopropyl ether.

25 **Example 183**

N-[4-[2-[(2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-3-methoxy-4-(2-quinolinylmethoxy)benzamide

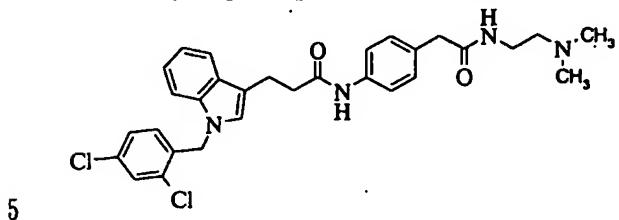


30 Melting point: 209-210 °C (decomposition).

Washing solvent: diisopropyl ether.

Example 184

3-[1-(2,4-Dichlorobenzyl)-1H-indol-3-yl]-N-[4-[2-[(2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]propanamide



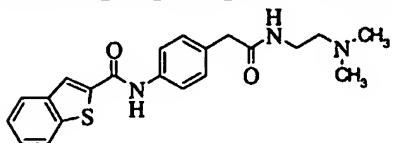
5

Melting point: 123-125 °C (decomposition).

Washing solvent: diisopropyl ether.

Example 185

10 N-[4-[2-[(2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-1-benzothiophen-2-carboxamide



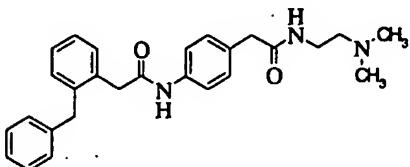
Melting point: 186-187 °C (decomposition).

Washing solvent: diisopropyl ether.

15

Example 186

2-(2-Benzylphenyl)-N-[4-[2-[(2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]acetamide

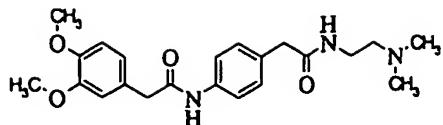


20 Melting point: 115-117 °C.

Washing solvent: diisopropyl ether.

Example 187

25 2-(3,4-dimethoxyphenyl)-N-[4-[2-[(2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]acetamide

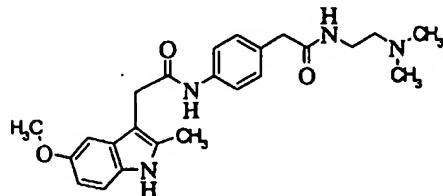


Melting point: 123-124 °C.

Recrystallization solvent: methanol - diisopropyl ether.

5 Example 188

N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-2-(5-methoxy-2-methyl-1H-indol-3-yl)acetamide

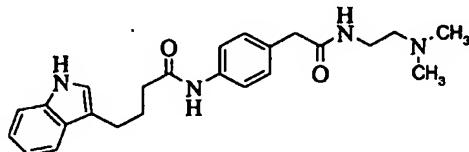


10 Melting point: 125-126 °C.

Recrystallization solvent: methanol - diisopropyl ether.

Example 189

N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-4-(1H-indol-3-yl)butanamide

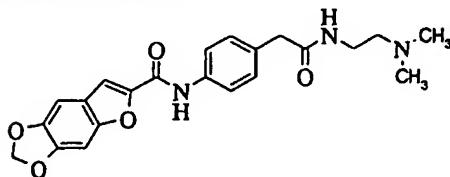


Melting point: 132-133 °C.

Washing solvent: diisopropyl ether.

20 Example 190

N-[4-[2-[[2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]furo[2,3-f][1,3]benzodioxol-6-carboxamide

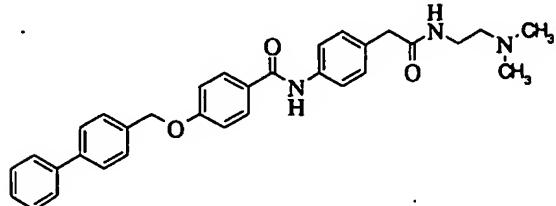


Melting point: 173-175 °C (decomposition).

Washing solvent: diisopropyl ether.

Example 191

5 4-((1,1'-Biphenyl)-4-ylmethoxy)-N-[4-[2-[(2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]benzamide



Melting point: 204-208 °C.

Washing solvent: diisopropyl ether.

10

Example 192

4-(Benzoylamino)-N-[4-[2-[(2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]benzamide

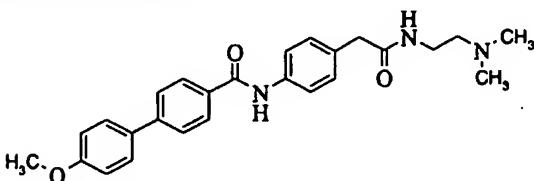


15 Melting point: 220-221 °C.

Washing solvent: diisopropyl ether.

Example 193

20 N-[4-[2-[(2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-4'-methoxy[1,1'-biphenyl]-4-carboxamide



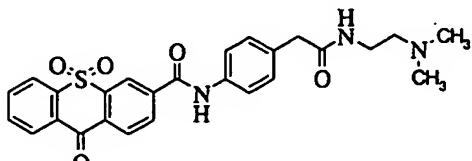
Melting point: 196-198 °C (decomposition).

Washing solvent: diisopropyl ether.

25

Example 194

N-[4-[2-[(2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-9,10,10-trioxo-9,10-dihydro-10λ⁶-thioxanten-3-carboxamide



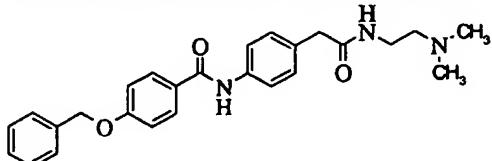
5

Melting point: 162-163 °C (decomposition).

Washing solvent: diisopropyl ether.

Example 195

10 4-(Benzylxy)-N-[4-[2-[(2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]benzamide

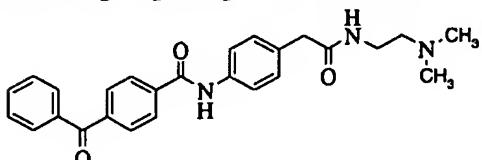


Melting point: 190-192 °C (decomposition).

15 Washing solvent: diisopropyl ether.

Example 196

4-Benzoyl-N-[4-[2-[(2-(dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]benzamide



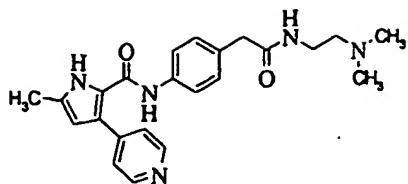
20

Melting point: 173-175 °C (decomposition).

Washing solvent: diisopropyl ether.

Example 197

25 N-[4-[2-[(2-(Dimethylamino)ethyl]amino]-2-oxoethyl]phenyl]-5-methyl-3-(4-pyridinyl)-1H-pyrole-2-carboxamide

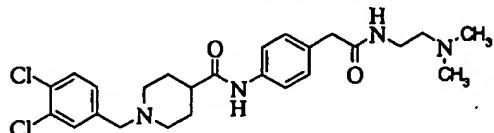


Melting point: 215-218 °C (decomposition).

Washing solvent: diisopropyl ether.

5 Example 198

1-(3,4-Dichlorobenzyl)-N-[4-{2-[(2-(dimethylamino)ethyl)amino]-2-oxoethyl}phenyl]-4-piperidinecarboxamide

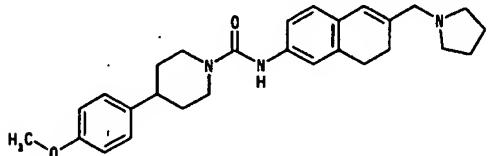


10 Melting point: 182-183 °C (decomposition).

Washing solvent: diisopropyl ether.

Example 199

15 4-(4-Methoxyphenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



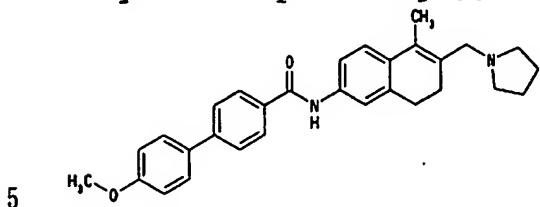
The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

20 ¹H-NMR (CDCl₃) δ: 1.61-1.91 (8H, m), 2.31 (2H, t, J=8.1 Hz), 2.54 (4H, m), 2.73-2.81 (3H, m), 2.98 (2H, t, J=7.8 Hz), 3.16 (2H, s), 3.79 (3H, s), 4.20 (2H, d, J=13.1 Hz), 6.31 (1H, s), 6.36 (1H, s), 6.86 (2H, d, J=8.6 Hz), 7.06-7.20 (5H, m).

25 Melting point: 175-176 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 200

4'-Methoxy-N-[6-(1-pyrrolidinylmethyl)-5-methyl-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained by carrying out the same operation as in Example 1, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

10 ¹H-NMR (CDCl₃) δ : 1.78 (4H, m), 2.10 (3H, s), 2.37 (2H, t, J=8.1 Hz), 2.53 (4H, m), 2.76 (2H, t, J=8.1 Hz), 3.28 (2H, s), 3.87 (3H, s), 7.01 (2H, d, J=8.6 Hz), 7.27 (1H, d, J=7.8 Hz), 7.46 (1H, d, J=7.8 Hz), 7.48 (1H, s), 7.57 (2H, d, J=8.6 Hz), 7.66 (2H, d, J=8.4 Hz), 7.81 (1H, s), 7.92 (2H, d, J=8.4 Hz).

15 Elemental analysis for C₃₀H₃₂N₂O₂

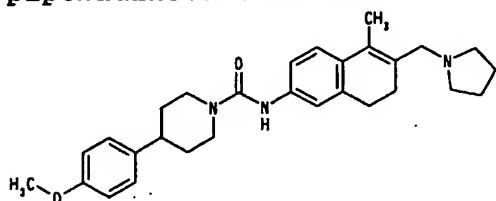
Calcd. : C, 79.61; H, 7.13; N, 6.19

Found : C, 79.35; H, 7.28; N, 6.24

20 Melting point: 179-180 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 201

4-(4-Methoxyphenyl)-N-[6-(1-pyrrolidinylmethyl)-5-methyl-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



The titled compound was obtained by carrying out the same operation as in Example 99, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine

obtained in Reference Example 69.

¹H-NMR (CDCl₃) δ : 1.67 (2H, dd, J=13.4, 4.0 Hz), 1.78 (4H, m), 1.89 (2H, d, J=11.4 Hz), 2.07 (3H, s), 2.34 (2H, t, J=7.5 Hz), 2.52 (4H, m), 2.68-2.73 (3H, m), 2.98 (2H, t, J=7.5 Hz), 5 3.26 (2H, s), 3.80 (3H, s), 4.20 (2H, d, J=13.4 Hz), 6.36 (1H, s), 6.86 (2H, d, J=8.4 Hz), 7.12-7.20 (5H, m).

Elemental analysis for C₂₈H₃₇N₃O₂

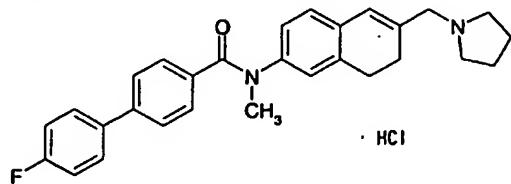
Calcd. : C, 75.13; H, 8.33; N, 9.39

Found : C, 74.96; H, 8.14; N, 9.10

10 Melting point: 163-164 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 202

4'-Fluoro-N-methyl-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide hydrochloride



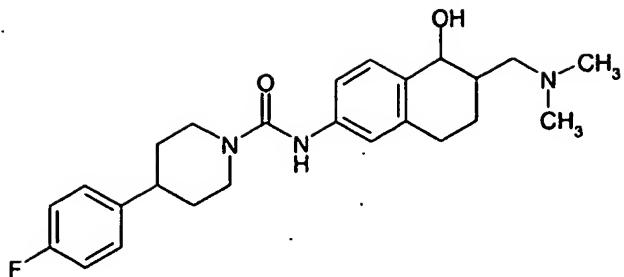
The titled compound was obtained by carrying out the same operation as in Example 1, using N-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine hydrochloride obtained in Reference Example 95.

¹H-NMR (DMSO-d₆) δ : 1.92-1.98 (4H, m), 2.39 (2H, t, J=8.1 Hz), 2.73 (2H, t, J=8.1 Hz), 3.00 (2H, m), 3.35 (3H, m), 3.44 (2H, m), 3.83 (2H, d, J=5.6 Hz), 6.62 (1H, s), 6.92-7.01 (2H, m), 7.11 (1H, s), 7.26 (2H, dd, J=8.9, 5.6 Hz), 7.38 (2H, d, J=8.1 Hz), 7.55 (2H, d, J=8.1 Hz), 7.69 (2H, dd, J=8.9, 5.6 Hz), 10.60 (1H, brs).

FABMS(pos) 441.2 [M+H]⁺

30 Example 203

N-[6-[(Dimethylamino)methyl]-5-hydroxy-5,6,7,8-tetrahydro-2-naphthalenyl]-4-(4-fluorophenyl)-1-piperidinecarboxamide



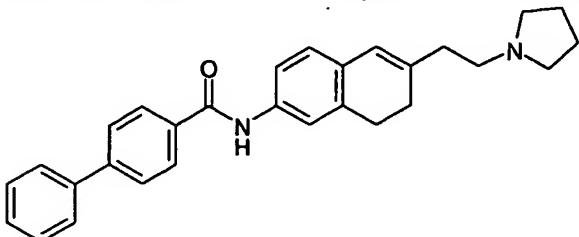
N, N-Dimethylmethylen ammonium chloride (638 mg, 6.82 mmol) was added to a mixed solution of 4-(4-fluorophenyl)-N-(5-oxo-5,6,7,8-tetrahydro-2-naphthalenyl)-1-piperidinecarboxamide (1.00 g, 2.73 mmol) obtained in Reference Example 97 in tetrahydrofuran (10 ml) and acetonitrile (10 ml), which was stirred at room temperature for 1 day. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting oily substance was dissolved in methanol (15 ml). Sodium borohydride (103 mg, 2.73 mmol) was added to the solution under ice-cooling, which was stirred for 1 hour. Then, the solvent was distilled out under reduced pressure. 1N Hydrochloric acid was added to the residue, which was washed with ethyl acetate. 4N Sodium hydroxide was added to the water layer to make it alkaline. The reaction mixture was extracted with ethyl acetate, which was washed with saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was purified by aluminum B column chromatography (development solvent; ethyl acetate), powdered with hexane, to give the titled compound (231 mg).

Melting point: 160-163 °C (crystallization solvent: ethyl acetate - n-hexane)

FAB(pos) 426.3 [M+H]+

Example 204

N-[6-[2-(1-Pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



5 Concentrated hydrochloric acid (2 ml) was added to N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl]acetamide (98.0 mg, 0.345 mmol) obtained in Reference Example 103, which was stirred at 100 °C for 16 hours. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. WSC (62.5 mg, 0.326 mmol) was added to a dimethylformamide solution (1.5ml) of the resulting oily substance (79.0 mg, 0.326 mmol), [1,1'-biphenyl]-4-carboxylic acid (64.6 mg, 0.326 mmol) and DMAP (39.8 mg, 0.326 mmol) under ice-cooling, which was stirred at room temperature for 1 day. Ethyl acetate was added to the reaction mixture, washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, then the solvent was distilled out under reduced pressure.

10 The resulting residue was purified by aluminum column chromatography (development solvent; ethyl acetate), powdered with ethyl acetate and isopropyl ether (1:5), to give the titled compound (36.8 mg).

15 ¹H NMR (DMSO-d₆) δ : 1.67 (4H, m), 2.23 (2H, m), 2.34 (2H, m), 2.46 (4H, m), 2.57 (2H, m), 2.75 (2H, m), 6.24 (1H, s), 6.98 (1H, d, J = 8.1 Hz), 7.40-7.59 (5H, m), 7.76 (2H, d, J = 7.5 Hz), 7.82 (2H, d, J=8.4 Hz), 8.05 (2H, d, J = 8.4

Hz), 10.19 (1H, s).

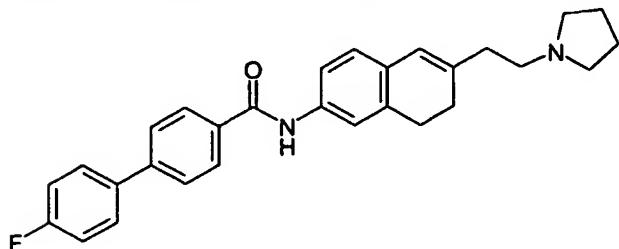
Melting point: 184-186 °C (crystallization solvent: ethyl acetate - isopropyl ether)

FAB(pos). 423.2 [M+H]⁺

5

Example 205

4'-Fluoro-N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



10 Concentrated hydrochloric acid (2 ml) was added to N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl]acetamide (98.0 mg, 0.345 mmol) obtained in Reference Example 103, which was stirred at 100°C for 16 hours. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. WSC (62.5 mg, 0.326 mmol) was added to a dimethylformamide solution (1.5 ml) of the resulting oily substance (79.0 mg, 0.326 mmol), 4'-fluoro-[1,1'-biphenyl]-4-carboxylic acid (64.6 mg, 0.326 mmol) and DMAP (39.8 mg, 0.326 mmol) under ice-cooling, which was stirred at room temperature for 1 day.

15 Ethyl acetate was added to the reaction mixture, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and the solvent was distilled out under reduced pressure. The resulting residue was purified by aluminum column chromatography (development solvent; ethyl acetate), powdered with ethyl acetate -

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isopropyl ether (1:5), to give the titled compound (75.1 mg).

¹H NMR (DMSO-d₆) δ : 1.68 (4H, m), 2.23 (2H, m), 2.35 (2H, m), 2.50 (4H, m), 2.59 (2H, m), 2.75 (2H, m), 6.24 (1H, s), 5 6.98 (1H, d, J = 8.1 Hz), 7.34 (2H, m), 7.56 (2H, m), 7.81 (4H, m), 8.04 (2H, d, J = 8.4 Hz), 10.19 (1H, s).

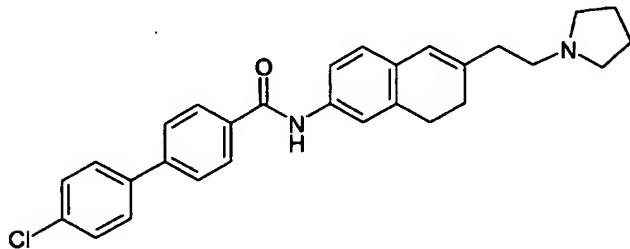
Melting point: 187-189°C (crystallization solvent: ethyl acetate - isopropyl ether)

FAB (pos) 441.3 [M+H]⁺

10

Example 206

4'-Chloro-N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



15 Concentrated hydrochloric acid (2 ml) was added to N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-naphthalenyl]acetamide (98.0 mg, 0.345 mmol) obtained in Reference Example 103, which was stirred at 100°C for 16 hours. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. WSC (62.5 mg, 0.326 mmol) was added to a dimethylformamide solution (1.5 ml) of the resulting oily substance (79.0 mg, 0.326 mmol), 4'-chloro-[1,1'-biphenyl]-4-carboxylic acid (64.6 mg, 0.326 mmol) and DMAP (39.8 mg, 0.326 mmol) under ice-cooling, which was stirred at room temperature for 1 day.

20 Ethyl acetate was added to the reaction mixture, which was washed with aqueous potassium carbonate solution and

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saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was purified by aluminum column chromatography
 5 (development solvent ; ethyl acetate), powdered with ethyl acetate - isopropyl ether (1:5), to give the titled compound (78.4 mg).

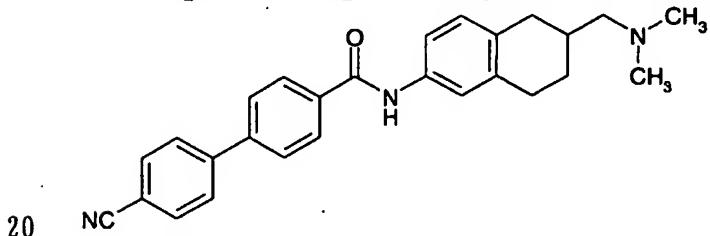
¹H NMR (DMSO-d₆) δ : 1.67 (4H, m), 2.23 (2H, m), 2.34 (2H, m), 2.45 (4H, m), 2.57 (2H, m), 2.75 (2H, m), 6.24 (1H, s),
 10 6.98 (1H, d, J = 8.1 Hz), 7.55 (4H, m), 7.80 (2H, d, J=8.4 Hz), 7.84 (2H, d, J=8.4 Hz), 8.05 (2H, d, J = 8.7 Hz), 10.20 (1H, s).

Melting point: 207-209°C (crystallization solvent: ethyl acetate - isopropyl ether)

15 FAB (pos) 457.2 [M+H]⁺

Example 207

4'-Cyano-N-[6-[(dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



20 The titled compound was obtained by carrying out the same operation as in Example 1, using N-[(6-amino-1,2,3,4-tetrahydro-2-naphthalenyl)methyl]-N,N-dimethylamine and 4'-cyano-[1,1'-biphenyl]-4-carboxylic acid.

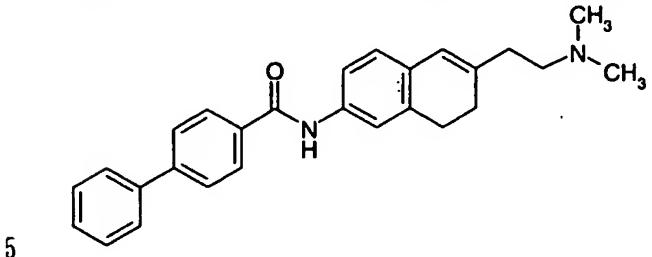
25 ¹H NMR (CDCl₃) δ : 1.42 (1H, m), 1.95 (2H, m), 2.26 (6H, s), 2.24-2.46 (3H, m), 2.84-2.95 (3H, m), 7.10 (1H, d, J=8.4 Hz), 7.30 (1H, m), 7.46 (1H, s), 7.74 (7H, m), 7.98 (2H, d, J=8.4 Hz).

30 Melting point: 183-185°C (crystallization solvent: ethyl acetate - isopropyl ether)

FAB (pos) 410.2 [M+H]⁺

Example 208

N-[6-[2-(Dimethylamino)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



Concentrated hydrochloric acid (1.5 ml) was added to N-[6-[2-(dimethylamino)ethyl]-7,8-dihydro-2-naphthalenyl]acetamide (57.5 mg, 0.223 mmol) obtained in Reference Example 104, which was stirred at 100°C for 1 hour. The solvent was distilled out under reduced pressure. Ethyl acetate was added to the residue, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. WSC (29.2 mg, 0.139 mmol) was added to a dimethylformamide solution (0.7 ml) of the resulting oily substance (30 mg, 0.139 mmol), [1,1'-biphenyl]-4-carboxylic acid (30.2 mg, 0.139 mmol) and DMAP (16.9 mg, 0.139 mmol) under ice-cooling, which was stirred at room temperature for 16 hours. Ethyl acetate was added to the reaction mixture, which was washed with aqueous potassium carbonate solution and saturated aqueous sodium chloride solution, dried over anhydrous sodium sulfate, and then the solvent was distilled out under reduced pressure. The resulting residue was purified by aluminum column chromatography (development solvent; ethyl acetate), powdered with ethyl acetate - isopropyl ether (1:5), to give the titled compound (12.4 mg).

¹H NMR (CDCl₃) δ: 2.29 (8H, m), 2.41 (2H, m), 2.46 (2H, m), 2.84 (2H, t, J = 8.1 Hz), 6.24 (1H, s), 6.98 (1H, d, J = 8.4 Hz), 7.34 (1H, m), 7.41 (1H, d, J = 6.9 Hz), 7.46 (3H,

10

15

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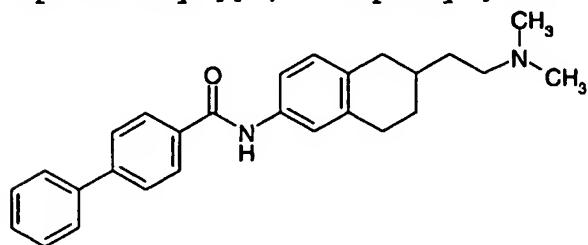
m), 7.63 (2H, d, J = 7.2 Hz), 7.71 (2H, d, J = 8.4 Hz), 7.77 (1H, br), 7.94 (2H, d, J = 8.4 Hz).

Melting point: 148-150°C (crystallization solvent: ethyl acetate - isopropyl ether)

5 FAB (pos) 397.2 [M+H]⁺

Example 209

N-[6-[2-(Dimethylamino)ethyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



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A methanol solution (5 ml) of N-[6-[2-(dimethylamino)ethyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide (20 mg, 0.050 mmol) obtained in Example 208 and palladium carbon (10 mg) was stirred under hydrogen atmosphere for 4 hours. After a catalyst was filtered off, the filtrate was concentrated under reduced pressure. The resulting residue was purified by aluminum B column chromatography (development solvent: ethyl acetate), powdered with ethyl acetate - hexane (1:3), to give the titled compound (4.0 mg).

15 ¹H NMR (CDCl₃) δ: 1.60 (4H, m), 1.92 (1H, m), 2.26 (6H, s), 2.42 (3H, m), 2.84 (3H, m), 7.06 (1H, d, J=8.1Hz), 7.32 (1H, m), 7.46 (4H, m), 7.63 (2H, d, J=6.9Hz), 7.72 (3H, m), 7.94 (2H, d, J=8.1Hz).

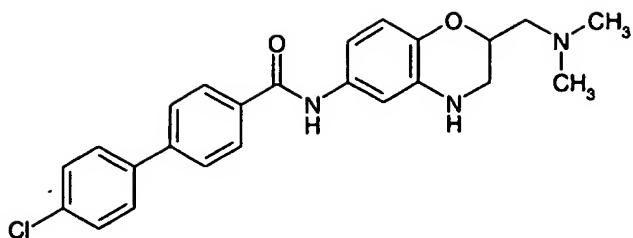
20 Melting point: 112-114°C (crystallization solvent: ethyl acetate - isopropyl ether)

25 FAB(pos) 399.2 [M+H]⁺

Example 210

30 4'-Chloro-N-[2-[(dimethylamino)methyl]-3,4-dihydro-2H-1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide

289



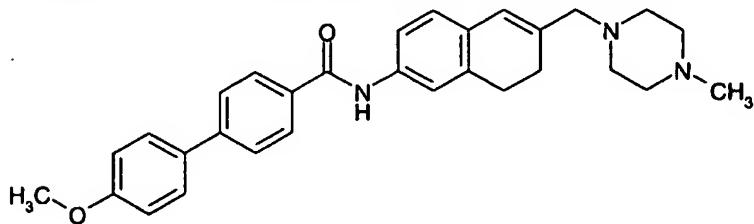
The titled compound was obtained as white powders by the same method as in Example 1, using 6-amino-2-(dimethylamino)methyl-1,4-benzoxazin obtained in Reference Example 105.

¹H-NMR (CDCl₃) δ : 2.33 (6H, s), 2.44-2.65 (2H, m), 3.15-3.21 (1H, m), 3.41-3.46 (1H, m), 3.87 (1H, brs), 4.24-4.26 (1H, m), 6.61 (1H, dd, J=2.5, 8.6 Hz), 6.81 (1H, d, J=8.6 Hz), 7.28 (1H, d, J=2.5 Hz), 7.43 (2H, d, J=6.5 Hz), 7.54 (2H, d, J=6.5 Hz), 7.64 (2H, d, J=8.4 Hz), 7.71 (1H, s), 7.90 (2H, d, J=8.4 Hz).

Melting point: 227-230 °C (crystallization solvent: diisopropyl ether)

15 Example 211

4'-Methoxy-N-[6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



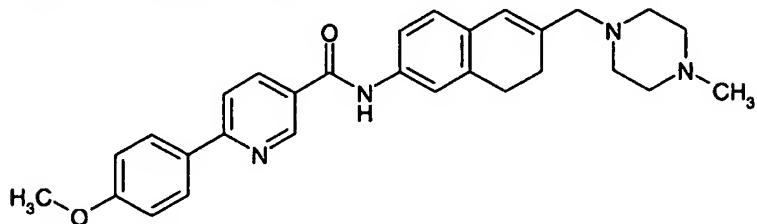
The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 106.

¹H NMR (CDCl₃) δ : 2.31 (3H, s), 2.33 (2H, t, J=8.1 Hz), 2.49 (8H, bs), 2.84 (2H, t, J=8.1 Hz), 3.07 (2H, s), 3.87 (3H, s), 6.36 (1H, s), 7.00-7.03 (3H, m), 7.36 (1H, d, J=8.1 Hz), 7.51 (1H, s), 7.58 (2H, d, J=8.4 Hz), 7.67 (2H, d, J=8.4 Hz), 7.78 (1H, s), 7.91 (2H, d, J=8.4 Hz).

Melting point: 208-210 °C (crystallization solvent: ethyl acetate)

Example 212

5 6-(4-Methoxyphenyl)-N-[6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]nicotinamide



The titled compound was obtained as colorless powders
 10 by carrying out the same operation as in Example 1, using
 6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 106.

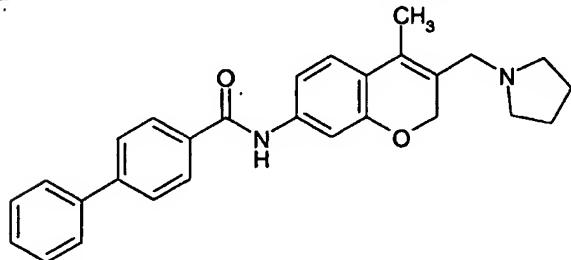
¹H NMR (CDCl₃) δ: 2.30 (3H, s), 2.33 (2H, t, J=8.1 Hz), 2.47 (8H, bs), 2.84 (2H, t, J=8.1 Hz), 3.07 (2H, s), 3.89 (3H, s), 6.36 (1H, s), 7.01-7.04 (3H, m), 7.37 (1H, d, J=8.1 Hz), 7.49 (1H, s), 7.78-7.81 (2H, m), 8.03 (2H, d, J=8.4 Hz), 8.21 (1H, dd, J=2.1 Hz, 8.7 Hz), 9.09 (1H, s).

Melting point: 235-237 °C (crystallization solvent: ethyl acetate)

20

Example 213

N-[4-Methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide



25 The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using

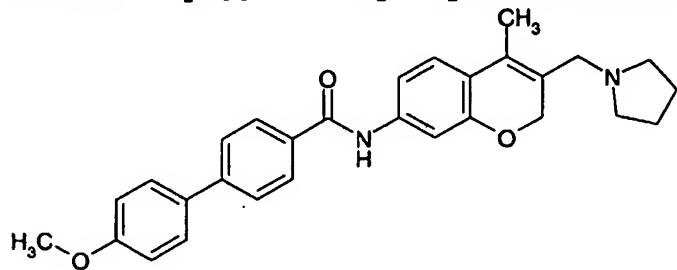
4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine
obtained in Reference Example 107.

¹H NMR (CDCl₃) δ : 1.77 (4H, s), 2.05 (3H, s), 2.51 (4H, s),
3.25 (2H, s), 4.74 (2H, s), 7.14-7.50 (6H, m), 7.63 (2H,
5 d, J=7.2 Hz), 7.71 (2H, d, J=8.4 Hz), 7.79 (1H, s), 7.94
(2H, d, J=8.4 Hz).

Melting point: 176-178 °C (crystallization solvent:
ethyl acetate - diisopropyl ether)

10 Example 214

4'-Methoxy-N-[4-methyl-3-(1-pyrrolidinylmethyl)-2H-
chromen-7-yl][1,1'-biphenyl]-4-carboxamide



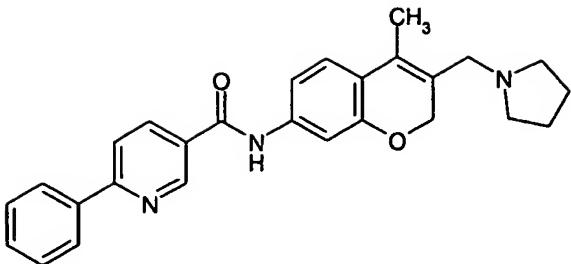
15 The titled compound was obtained as colorless powders
by carrying out the same operation as in Example 1, using
4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine
obtained in Reference Example 107.

¹H NMR (CDCl₃) δ : 1.77 (4H, s), 2.05 (3H, s), 2.51 (4H, s),
3.25 (2H, s), 3.87 (3H, s), 4.74 (2H, s), 7.01 (2H, d, J=8.7
20 Hz), 7.14-7.31 (3H, m), 7.57 (2H, d, J=8.7 Hz), 7.66 (2H,
d, J=8.4 Hz), 7.89 (1H, s), 7.91 (2H, d, J=8.4 Hz).

Melting point: 195-197 °C (crystallization solvent:
ethyl acetate - diisopropyl ether)

25 Example 215

N-[4-Methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]-
6-phenylnicotinamide



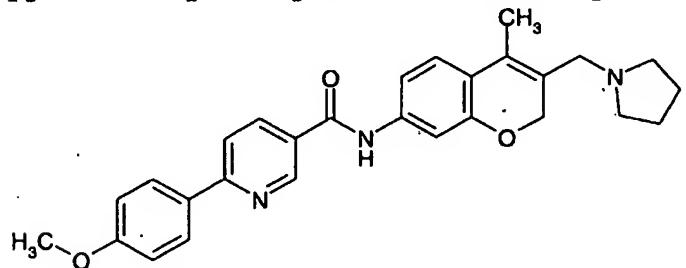
The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 107.

¹H NMR (CDCl₃) δ : 1.77 (4H, s), 2.05 (3H, s), 2.51 (4H, s), 3.25 (2H, s), 4.74 (2H, s), 7.14-7.28 (3H, m), 7.47-7.54 (3H, m), 7.81-7.87 (2H, m), 8.06 (2H, d, J=8.4 Hz), 8.27 (1H, d, J=8.4 Hz), 9.13 (1H, s).

Melting point: 192-193 °C (crystallization solvent: ethyl acetate)

Example 216

15 6-(4-Methoxyphenyl)-N-[4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]nicotinamide



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 107.

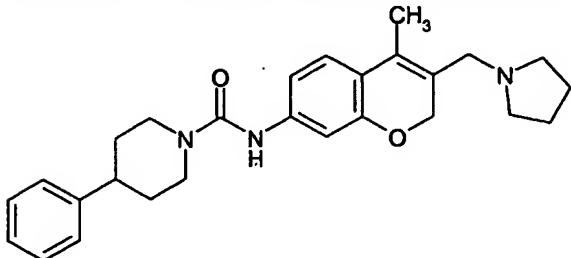
¹H NMR (CDCl₃) δ : 1.77 (4H, s), 2.05 (3H, s), 2.51 (4H, s), 3.25 (2H, s), 3.89 (3H, s), 4.74 (2H, s), 7.03 (2H, d, J=8.7 Hz), 7.14-7.26 (3H, m), 7.75-7.81 (2H, m), 8.03 (2H, d, J=8.7 Hz), 8.21 (1H, d, J=6.6 Hz), 9.09 (1H, s).

Melting point: 201-203 °C (crystallization solvent: ethyl

acetate)

Example 217

N-[4-Methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]-
5 4-phenyl-1-piperidinecarboxamide



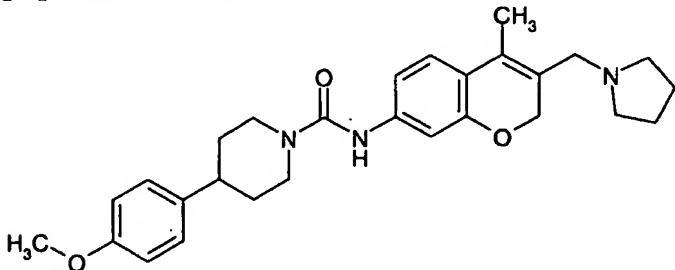
The titled compound was obtained as colorless powders by carrying out the same operation as in Example 99, using
4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine
10 obtained in Reference Example 107.

¹H NMR (CDCl₃) δ : 1.72-1.95 (8H, m), 2.03 (3H, s), 2.54 (4H, s), 2.63-2.76 (1H, m), 2.95-3.00 (2H, m), 3.27 (2H, s), 4.19-4.23 (2H, m), 4.70 (2H, s), 6.39 (1H, s), 6.83 (1H, s), 7.01-7.32 (7H, m).

15 Melting point: 125-127 °C (crystallization solvent:
ethyl acetate - diisopropyl ether)

Example 218

4-(4-Methoxyphenyl)-N-[4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]-1-piperidinecarboxamide



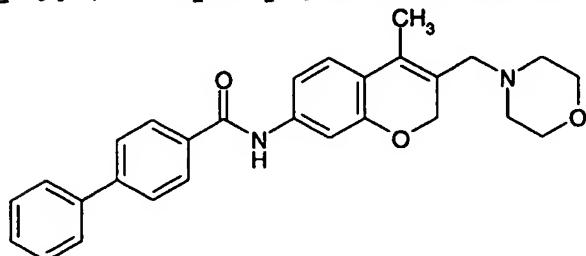
The titled compound was obtained as colorless powders by carrying out the same operation as in Example 99, using
25 4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine

obtained in Reference Example 107.

¹H NMR (CDCl₃) δ : 1.63-1.91 (8H, m), 2.02 (3H, s), 2.49 (4H, s), 2.61-2.71 (1H, m), 2.93-3.01 (2H, m), 3.23 (2H, s), 3.79 (3H, s), 4.16-4.21 (2H, m), 4.69 (2H, s), 6.34 (1H, s),
5 6.82-6.91 (3H, m), 6.99-7.02 (1H, m), 7.10-7.15 (3H, m).
Melting point: 144-146 °C (crystallization solvent: ethyl acetate - n-hexane)

Example 219

10 N-[4-Methyl-3-(4-morpholinylmethyl)-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide

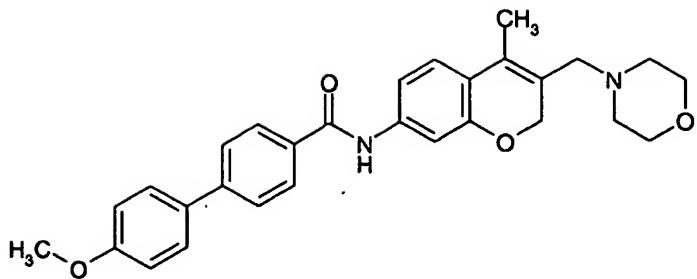


The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using
15 4-methyl-3-(4-morpholinylmethyl)-2H-chromen-7-amine obtained in Reference Example 108.

¹H NMR (DMSO-d₆) δ : 2.01 (3H, s), 2.37 (4H, s), 3.32 (2H, s), 3.57 (4H, s), 4.63 (2H, s), 7.23 (1H, d, J=8.1 Hz), 7.38-7.54 (5H, m), 7.76 (2H, d, J=7.5 Hz), 7.84 (2H, d, J=8.1 Hz), 8.04 (2H, d, J=8.1 Hz), 10.27 (1H, s).
20 Melting point: 162-164 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 220

25 4'-Methoxy-N-[4-methyl-3-(4-morpholinylmethyl)-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide



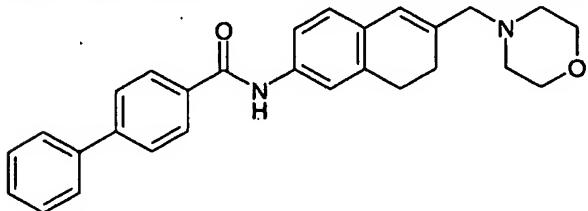
The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 4-methyl-3-(4-morpholinylmethyl)-2H-chromen-7-amine obtained in Reference Example 108.

¹H NMR (DMSO-d₆) δ : 2.00 (3H, s), 2.37 (4H, s), 3.11 (2H, s), 3.57 (4H, s), 3.82 (3H, s), 4.63 (2H, s), 7.07 (2H, d, J=8.7 Hz), 7.23 (1H, d, J=8.1 Hz), 7.38-7.40 (2H, m), 7.72 (2H, d, J=8.7 Hz), 7.79 (2H, d, J=8.4 Hz), 8.01 (2H, d, J=8.4 Hz), 10.23 (1H, s).

Melting point: 198-200 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 221

N-[6-(4-Morpholinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-(4-morpholinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 109.

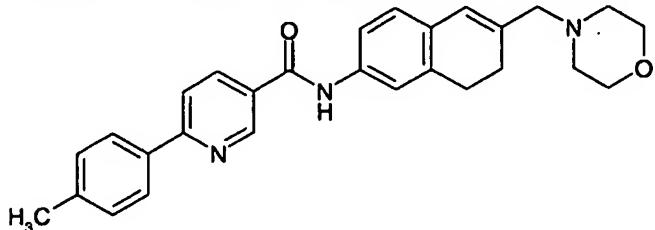
¹H-NMR (CDCl₃) δ : 2.34 (2H, t, J=8.4 Hz), 2.45 (4H, m), 2.85 (2H, t, J=8.4 Hz), 3.06 (2H, s), 3.73 (4H, t, J=4.7 Hz), 6.36 (1H, s), 7.02 (1H, d, J=8.1 Hz), 7.36-7.78 (10H, m), 7.93 (2H, d, J=8.1 Hz).

Melting point: 180-181 °C (crystallization solvent:

ethyl acetate - diisopropyl ether)

Example 222

6-(4-Methylphenyl)-N-[6-(4-morpholinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide



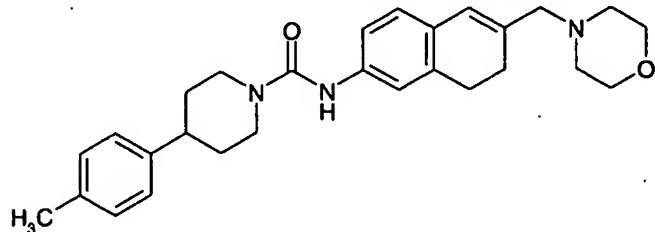
The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-(4-morpholinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 109.

¹H-NMR (CDCl₃) δ : 2.39 (2H, t, J=8.4 Hz), 2.43 (7H, m), 2.85 (2H, t, J=8.4 Hz), 3.06 (2H, s), 3.73 (4H, t, J=4.5 Hz), 6.36 (1H, s), 7.03 (1H, d, J=8.1 Hz), 7.30-7.38 (3H, m), 7.50 (1H, s), 7.76 (1H, s), 7.84 (1H, d, J=8.1 Hz), 7.97 (2H, d, J=8.1 Hz), 8.24 (1H, dd, J=8.4, 2.3 Hz), 9.12 (1H, s).

Melting point: 233-234 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 223

4-(4-Methylphenyl)-N-[6-(4-morpholinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



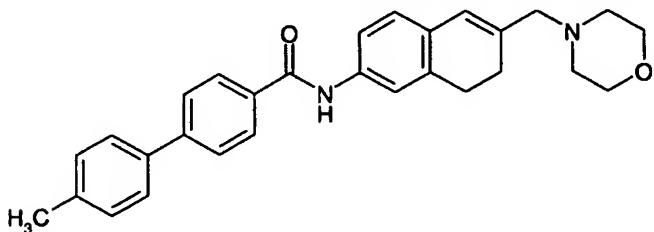
The titled compound was obtained as colorless powders by carrying out the same operation as in Example 99, using 6-(4-morpholinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 109.

¹H-NMR (CDCl₃) δ : 1.65-1.75 (4H, m), 1.90 (2H, m), 2.27-2.43 (7H, m), 2.72 (1H, m), 2.79 (2H, t, J=7.5 Hz), 2.93-3.04 (4H, m), 3.72 (4H, m), 4.20 (2H, d, J=11.7 Hz), 6.31 (1H, s), 6.39 (1H, s), 6.92 (1H, d, J=8.1 Hz), 7.05-7.26 (6H, m).

Melting point: 231-214 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 224

10 4'-Methyl-N-[6-(4-morpholinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



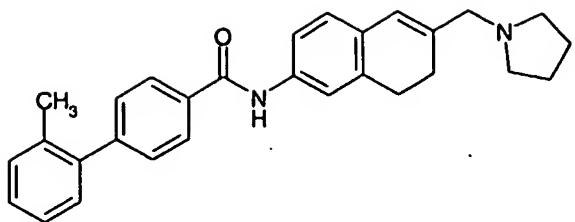
The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using
15 6-(4-morpholinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 109.

¹H-NMR (CDCl₃) δ : 2.33 (2H, t, J=8.1 Hz), 2.42-2.44 (7H, m), 2.84 (2H, t, J=8.1 Hz), 3.06 (2H, s), 3.72 (4H, t, J=4.2 Hz), 6.36 (1H, s), 7.01 (1H, d, J=8.1 Hz), 7.25-7.29 (2H, m), 7.37 (1H, d, J=8.1 Hz), 7.51-7.54 (3H, m), 7.68 (2H, d, J=8.1 Hz), 7.85 (1H, s), 7.92 (2H, d, J=8.1 Hz).

Melting point: 196-197 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

25 Example 225

2'-Methyl-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

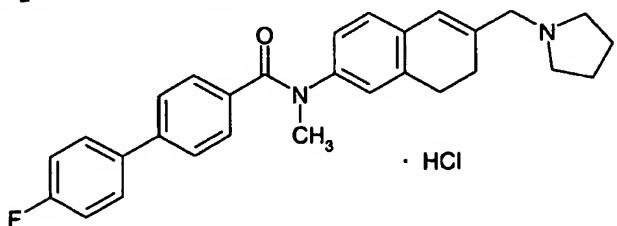


The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine 5 obtained in Reference Example 54.

Melting point: 177-178 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 226

10 4'-Fluoro-N-methyl-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide Hydrochloride



15 N-Methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine dihydrochloride (315 mg, 1.0 mmol) obtained in Reference Example 113 was dissolved in N,N-dimethylformamide (25 ml). 4-Bromobenzoic acid (402 mg, 2.0 mmol), WSC (383 mg, 2.0 mmol), HOEt (270 mg, 2.0 mmol) and DMAP (244 mg, 2.0 mmol) were added to the solution, 20 which was stirred at room temperature for 16 hours. Ethyl acetate and water were added to the reaction mixture, and extraction was conducted. The ethyl acetate layer was concentrated under reduced pressure. The residue was purified by aluminum column chromatography (development 25 solvent: ethyl acetate : n-hexane = 33:67). The eluate was concentrated under reduced pressure, which was dissolved in dimethoxyethane - tetrahydrofuran (10:1, 5.5 ml).

4-Fluorophenylboric acid (73 mg, 0.52 mmol), tetrakis(triphenylphosphine)palladium complex (15 mg, 0.013 mmol) and 2N aqueous sodium carbonate solution (0.433 ml) were added to the solution, which was refluxed 5 with heating under nitrogen atmosphere at 90°C for 5.5 hours. The reaction mixture was poured into cold water, and extraction was conducted using ethyl acetate. The ethyl acetate layer was concentrated, and the residue was purified by aluminum column chromatography (development 10 solvent; ethyl acetate). 4N Hydrogen chloride - ethyl acetate solution was added to the eluate, which was concentrated under reduced pressure. The resulting residue was recrystallized from methanol - ethyl acetate, to give the titled compound (108 mg).

15 $^1\text{H-NMR}$ (DMSO-d₆) δ : 1.92-1.98 (4H, m), 2.39 (2H, t, J=8.1 Hz), 2.73 (2H, t, J=8.1 Hz), 3.00 (2H, m), 3.35 (3H, m), 3.44 (2H, m), 3.83 (2H, d, J=5.6 Hz), 6.62 (1H, s), 6.92-7.01 (2H, m), 7.11 (1H, s), 7.26 (2H, dd, J=8.9, 5.6 Hz), 7.38 (2H, d, J=8.1 Hz), 7.55 (2H, d, J=8.1 Hz), 7.69 (2H, dd, J=8.9, 5.6 Hz), 10.60 (1H, brs.).

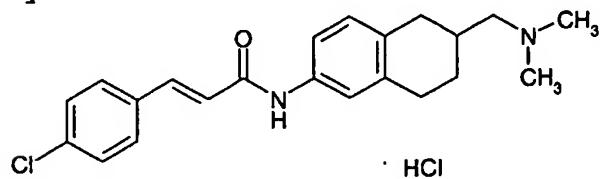
20 Melting point: 201-203 °C (crystallization solvent: methanol - diisopropyl ether)

FAB(pos) 441.2 [M+H]⁺

25 Example 227

(E)-3-(4-Chlorophenyl)-N-[6-[(dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl]-2-propenamide

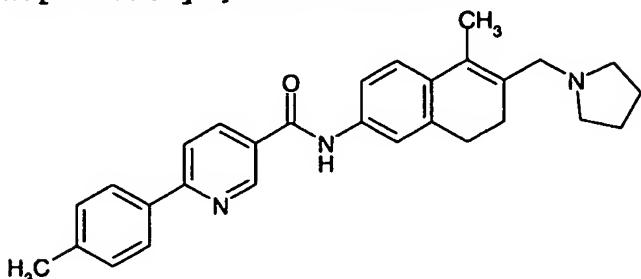
Hydrochloride



30 The titled compound was obtained as colorless powders by carrying out the same operation as in Example 4. Melting point: 243-245 °C (crystallization solvent: methanol - diisopropyl ether)

Example 228

6-(4-Methylphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

Melting point: 175-176 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

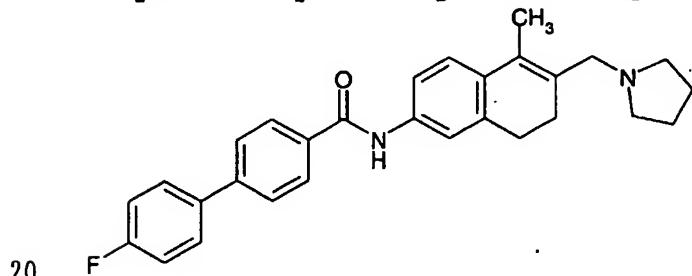
Elemental analysis for C₂₉H₃₀N₃O

Calcd.: C, 79.78; H, 6.93; N, 9.63

Found: C, 79.66; H, 6.97; N, 9.68

Example 229

4'-Fluoro-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

Melting point: 199-201 °C (crystallization solvent: ethyl

acetate - diisopropyl ether)

Elemental analysis for C₂₉H₃₀FN₂O

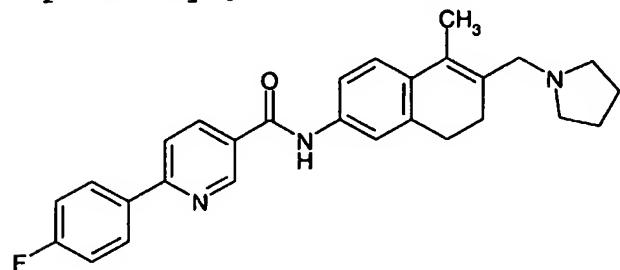
Calcd.: C, 79.06; H, 6.63; N, 6.36

Found: C, 79.01; H, 6.81; N, 6.45

5

Example 230

6-(4-Fluorophenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]nicotinamide



10

The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

15

Melting point: 204-205 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Elemental analysis for C₂₈H₂₈FN₂O

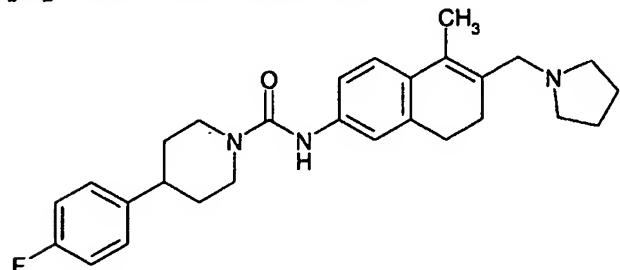
Calcd.: C, 76.17; H, 6.39; N, 9.52

Found: C, 76.03; H, 6.44; N, 9.62

20

Example 231

4-(4-Fluorophenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



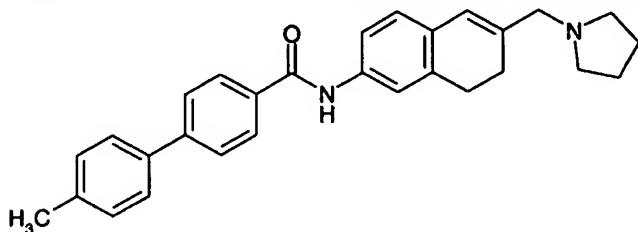
25

The titled compound was obtained as colorless powders by carrying out the same operation as in Example 99, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

5 Melting point: 172-173 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 232

10 4'-Methyl-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

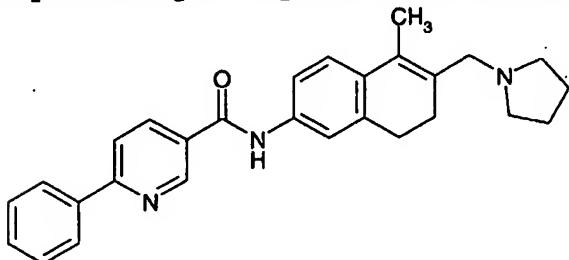


15 The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

Melting point: 176-177 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 233

20 N-[5-Methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-6-phenylnicotinamide



25 The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

Melting point: 178-179 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

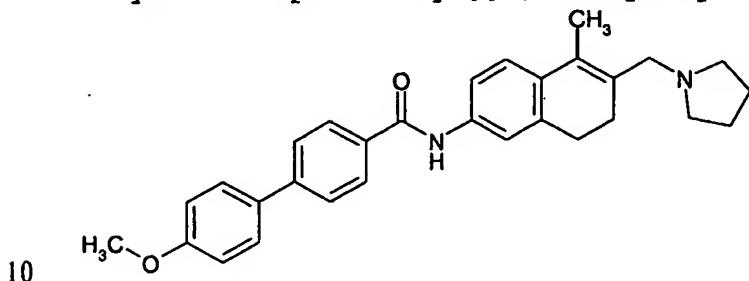
Elemental analysis for C₂₈H₂₉N₃O

Calcd.: C, 79.40; H, 6.90; N, 9.92

5 Found: C, 79.13; H, 6.82; N, 10.03

Example 234

4'-Methoxy-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



10

The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

15

¹H-NMR (CDCl₃) δ: 1.78 (4H, m), 2.10(3H, s), 2.37 (2H, t, J=8.1 Hz), 2.53 (4H, m), 2.76 (2H, t, J=8.1 Hz), 3.28(2H, s), 3.87 (3H, s), 7.01 (1H, d, J=8.6 Hz), 7.27 (2H, d, J=7.8 Hz), 7.46 (1H, d, J=7.8 Hz), 7.48 (1H, s), 7.57 (2H, d, J=8.6 Hz), 7.66 (2H, d, J=8.6 Hz), 7.81 (1H, s), 7.92 (2H, d, J=7.8 Hz).

20

Melting point: 179-180 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Elemental analysis for C₃₀H₃₂N₂O₂

Calcd.: C, 79.61; H, 7.13; N, 6.19

25

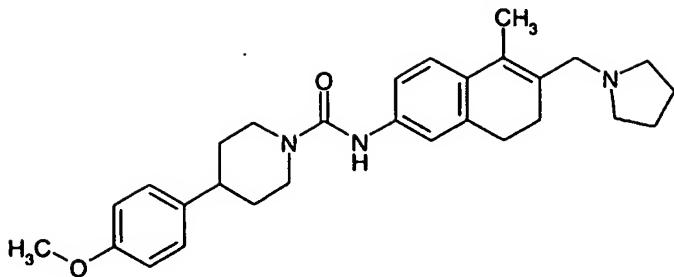
Found: C, 79.35; H, 7.28; N, 6.24

Example 235

4-(4-Methoxyphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide

30

304



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 99, using 5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

¹H-NMR (CDCl₃) δ : 1.67 (2H, dd, J=13.4, 4.0 Hz), 1.78 (4H, m), 1.89 (2H, d, J=11.4 Hz), 2.07 (3H, s), 2.34 (2H, t, J=7.5 Hz), 2.52 (4H, m), 2.68-2.73 (3H, m), 2.98 (2H, t, J=7.5 Hz), 3.26 (2H, s), 3.80 (3H, s), 4.20 (2H, d, J=13.4 Hz), 6.36 (1H, s), 6.86 (2H, d, J=8.4 Hz), 7.12-7.20 (5H, m).

Melting point: 163-164 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

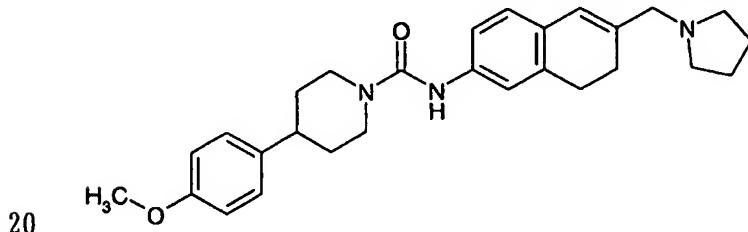
Elemental analysis for C₂₈H₃₁N₃O₂

Calcd. : C, 75.13; H, 8.33; N, 9.39

15 Found : C, 74.96; H, 8.14; N, 9.10

Example 236

4-(4-Methoxyphenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

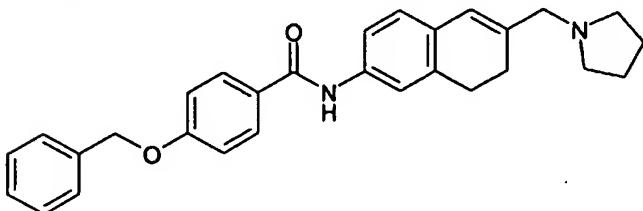
25 ¹H-NMR (CDCl₃) δ : 1.61-1.91 (8H, m), 2.31 (2H, t, J=8.1 Hz), 2.54 (4H, m), 2.73-2.81 (3H, m), 2.98 (2H, t, J=7.8 Hz),

3.16 (2H, s), 3.79 (3H, s), 4.20 (2H, d, J=13.1 Hz), 6.31 (1H, s), 6.36 (1H, s), 6.86 (2H, d, J=8.6 Hz), 7.06-7.20 (5H, m).

5 Melting point: 175-176 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 237

4-(Benzylloxy)-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]benzamide



10

The titled compound was obtained as colorless powders by carrying out the same operation as in Example 1, using 6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 54.

15 Melting point: 174-175 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Elemental analysis for C₂₈H₃₀N₂O₂

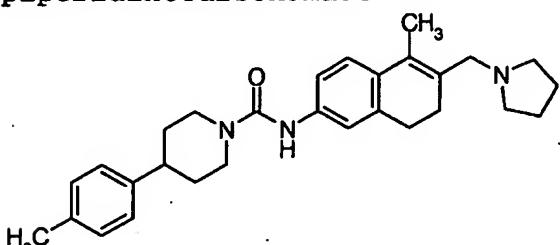
Calcd.: C, 78.84; H, 7.09; N, 6.87

Found: C, 79.06; H, 6.99; N, 6.41

20

Example 238

4-(4-Methylphenyl)-N-[5-methyl-6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



25

The titled compound was obtained by carrying out the same operation as in Example 99, using 5-methyl-6-(1-

pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenamine obtained in Reference Example 69.

¹H-NMR (CDCl₃) δ: 1.65-1.78 (6H, m), 1.90 (2H, d, J=12.9 Hz), 2.07 (3H, s), 2.33-2.37 (5H, m), 2.53 (4H, m), 5 2.68-2.74 (3H, m), 2.99 (2H, m), 3.27(2H,s), 4.21 (2H, d, J=13.2 Hz), 6.37 (1H, s), 7.09-7.21 (7H, m).

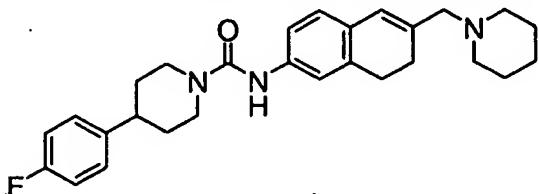
Melting point: 159-160 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

FAB(pos) 444.3 [M+H]⁺

10

Example 239

4-(4-Fluorophenyl)-N-[6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



15 The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenamine dihydrochloride obtained in Reference Example 114.

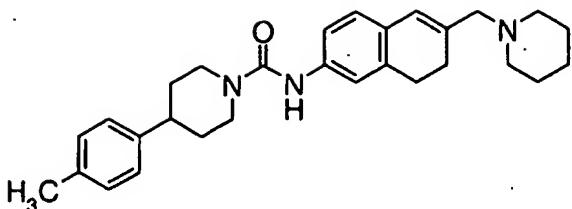
¹H-NMR (CDCl₃) δ: 1.43 (2H, m), 1.56-1.75 (6H, m), 1.89 (2H, d, J=12.3 Hz), 2.27-2.36 (6H, m), 2.70 (1H, m), 2.78 (2H, t, J=7.5 Hz), 2.88-3.00 (4H, m), 4.20 (2H, d, J=13.2 Hz), 6.29 (1H, s), 6.38 (1H, s), 6.91-7.08 (4H, m), 7.14-7.20(3H, m).

Melting point: 194 -195 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

25

Example 240

4-(4-Methylphenyl)-N-[6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



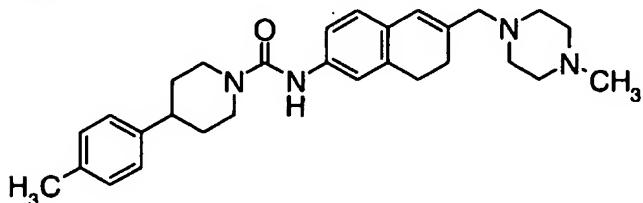
The titled compound was obtained by carrying out the same operation as in Example 99, using 6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenamine dihydrochloride obtained in Reference Example 114.

¹H-NMR (CDCl₃) δ : 1.43 (2H, m), 1.56-1.74 (6H, m), 1.90 (2H, d, J=12.0 Hz), 2.27-2.36 (9H, m), 2.69 (1H, m), 2.79 (2H, t, J=8.1 Hz), 2.94-3.01 (4H, m), 4.19 (2H, d, J=13.2 Hz), 6.29 (1H, s), 6.35 (1H, s), 6.93 (2H, d, J=8.1 Hz), 7.05-7.26 (5H, m).

Melting point: 209 -210 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

Example 241

15 4-(4-Methylphenyl)-N-[6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



The titled compound was obtained by carrying out the same operation as in Example 99, using 6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 106.

¹H NMR (CDCl₃) δ : 1.62-1.77 (2H, m), 1.90 (2H, d, J=12.0 Hz), 2.28 (2H, t, J=8.1 Hz), 2.29 (3H, s), 2.33 (3H, s), 2.46 (8H, bs), 2.64-2.73 (1H, m), 2.79 (2H, t, J=8.1 Hz), 2.96 (2H, d, J=10.5 Hz), 3.05 (2H, s), 4.19 (2H, d, J=13.5 Hz), 6.31 (1H, s), 6.34 (1H, s), 6.93 (1H, d, J=8.4 Hz), 7.04-7.16 (5H, m), 7.23 (1H, s).

Melting point: 214-216 °C (crystallization solvent:
tetrahydrofuran - n-hexane)

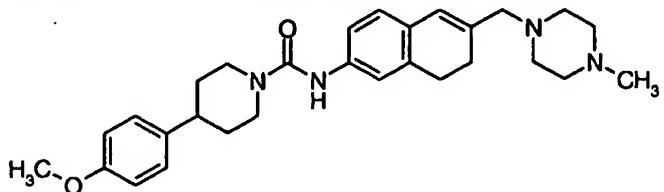
Elemental analysis for C₂₉H₃₈N₄O

Calcd.: C, 75.94; H, 8.35; N, 12.22.

5 Found: C, 75.67; H, 8.47; N, 12.27.

Example 242

4-(4-Methoxyphenyl)-N-[6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-
10 piperidinecarboxamide



The titled compound was obtained by carrying out the same operation as in Example 99, using 6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 106.

15 ¹H NMR (CDCl₃) δ: 1.68-1.76 (2H, m), 1.89 (2H, d, J=11.1 Hz), 2.29 (2H, t, J=8.1 Hz), 2.29 (3H, s), 2.46 (8H, bs), 2.64-2.71 (1H, m), 2.79 (2H, t, J=8.1 Hz), 2.82-3.03 (2H, m), 3.05 (2H, s), 3.80 (3H, s), 4.19 (2H, d, J=12.6 Hz), 20 6.31 (1H, s), 6.34 (1H, s), 6.87 (2H, d, J=8.7 Hz), 6.93 (1H, d, J=8.4 Hz), 7.06 (1H, dd, J=8.1, 2.1 Hz), 7.14 (2H, d, J=8.7 Hz), 7.23 (1H, s).

Melting point: 198-200 °C (crystallization solvent:
tetrahydrofuran - n-hexane)

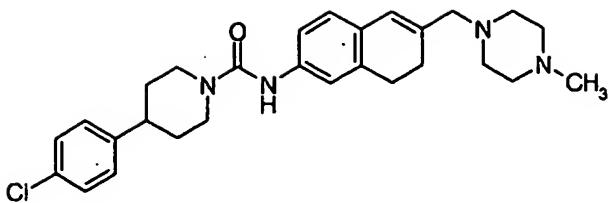
25 Elemental analysis for C₂₉H₃₈N₄O₂

Calcd.: C, 73.38; H, 8.07; N, 11.80.

Found: C, 73.04; H, 7.95; N, 11.67.

Example 243

30 4-(4-Chlorophenyl)-N-[6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



The titled compound was obtained by carrying out the same operation as in Example 99, using 6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine

5 obtained in Reference Example 106.

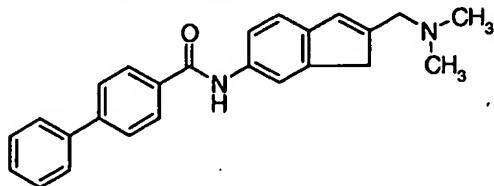
¹H NMR (CDCl₃) δ: 1.64-1.76 (2H, m), 1.90 (2H, d, J=11.1 Hz), 2.29 (2H, t, J=8.1 Hz), 2.29 (3H, s), 2.46 (8H, bs), 2.66-2.72 (1H, m), 2.79 (2H, t, J=8.1 Hz), 2.81-3.03 (2H, m), 3.05 (2H, s), 4.20 (2H, d, J=12.6 Hz), 6.31 (1H, s), 10 6.34 (1H, s), 6.93 (1H, d, J=7.8 Hz), 7.04-7.07 (1H, m), 7.14 (2H, d, J=8.4 Hz), 7.22 (1H, s), 7.28 (2H, d, J=8.4 Hz).

Melting point: 201-203 °C (crystallization solvent: tetrahydrofuran - n-hexane)

15

Example 244

N-[2-[(Dimethylamino)methyl]-1H-inden-6-yl][1,1'-biphenyl]-4-carboxamide



20 The titled compound was obtained by carrying out the same operation as in Example 1, using 2-[(dimethylamino)methyl]-1H-inden-6-amine obtained in Reference Example 116.

Elemental analysis for C₂₅H₂₄N₂O · 0.5H₂O

25 Calcd.: C, 79.55; H, 6.68; N, 7.42.

Found: C, 79.38; H, 6.76; N, 7.34.

Melting point: 187-189 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

FAB(pos) 369.2 [M+H]⁺

Example 245

5 N-[2-[(Dimethylamino)methyl]-1H-inden-6-yl]-4'-fluoro[1,1'-biphenyl]-4-carboxamide



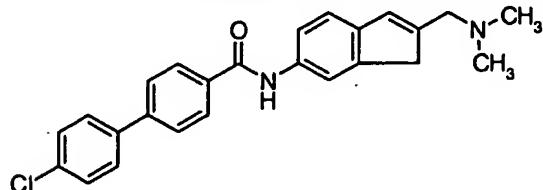
The titled compound was obtained by carrying out the same operation as in Example 1, using 2-[(dimethylamino)methyl]-1H-inden-6-amine obtained in
10 Reference Example 116.

Melting point: 209-211 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

FAB(pos) 387.2 [M+H]⁺

15 **Example 246**

4'-Chloro-N-[2-[(dimethylamino)methyl]-1H-inden-6-yl][1,1'-biphenyl]-4-carboxamide



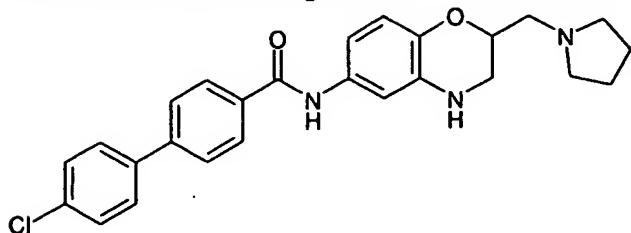
The titled compound was obtained by carrying out the same operation as in Example 1, using 2-[(dimethylamino)methyl]-1H-inden-6-amine obtained in
20 Reference Example 116.

Melting point: 218-220 °C (crystallization solvent: ethyl acetate - diisopropyl ether)

25 FAB(pos) 403.2 [M+H]⁺

Example 247

4'-Chloro-N-[2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-

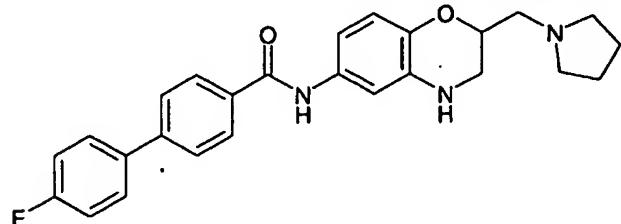
1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide

5 The titled compound was obtained by carrying out the same operation as in Example 1, using 6-amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine obtained in Reference Example 117.

10 $^1\text{H-NMR}$ (CDCl_3) δ : 1.70-1.90 (4H, m), 2.50-2.70 (4H, m), 2.73 (2H, d, $J=6.0\text{Hz}$), 3.18-3.24 (1H, m), 3.45-3.49 (1H, m), 3.87 (1H, brs), 4.26-4.28 (1H, m), 6.61 (1H, dd, $J=2.7, 8.4\text{ Hz}$), 6.80 (1H, d, $J=8.4\text{ Hz}$), 7.26 (1H, d, $J=2.7\text{ Hz}$), 7.44 (2H, d, $J=8.4\text{ Hz}$), 7.55 (2H, d, $J=8.4\text{ Hz}$), 7.64 (2H, d, $J=8.1\text{ Hz}$), 7.71 (1H, s), 7.91 (2H, d, $J=8.1\text{ Hz}$).

Melting point: 221-222 °C (crystallization solvent: diisopropyl ether)

15

Example 248**4'-Fluoro-N-[2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide**

20 The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 6-amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine obtained in Reference Example 117.

25 $^1\text{H-NMR}$ (CDCl_3) δ : 1.70-1.90 (4H, m), 2.50-2.70 (4H, m), 2.73 (2H, d, $J = 6.3\text{Hz}$), 3.18-3.24 (1H, m), 3.45-3.49 (1H, m), 3.88 (1H, brs), 4.24-4.30 (1H, m), 6.62 (1H, dd, $J=2.7, 8.4\text{ Hz}$), 6.80 (1H, d, $J=8.4\text{ Hz}$), 7.13-7.19 (2H, m), 7.26 (1H,

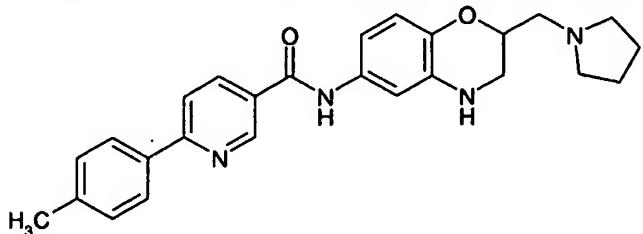
d, J=2.7 Hz), 7.56-7.60 (2H, m), 7.63 (2H, d, J=8.4 Hz), 7.71 (1H, s), 7.90 (2H, d, J=8.4 Hz).

Melting point: 204-206 °C (crystallization solvent: diisopropyl ether)

5

Example 249

6-(4-Methylphenyl)-N-[2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazin-6-yl]nicotinamide



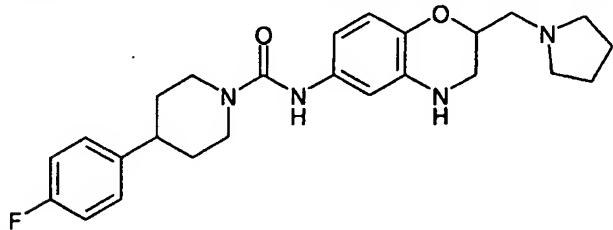
10 The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 6-amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine obtained in Reference Example 117.

15 $^1\text{H-NMR}$ (CDCl_3) δ : 1.70-1.85 (4H, m), 2.43 (3H, s), 2.50-2.70 (4H, m), 2.74 (2H, d, J=6.3Hz), 3.19-3.25 (1H, m), 3.45-3.49 (1H, m), 3.90 (1H, brs), 4.27-4.29 (1H, m), 6.63 (1H, dd, J=2.4, 8.7 Hz), 6.81 (1H, d, J=8.7 Hz), 7.26 (1H, d, J=2.7 Hz), 7.31 (2H, d, J=8.1 Hz), 7.67 (1H, s), 7.81 (1H, d, J=8.1 Hz), 7.93 (2H, d, J=7.8Hz), 8.21 (1H, dd, J=2.4, 8.4 Hz), 20 9.09 (1H, d, J=2.4 Hz).

Melting point: 207-208 °C (crystallization solvent: diisopropyl ether)

Example 250

25 4-(4-Fluorophenyl)-N-[2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazin-6-yl]-1-piperidinecarboxamide



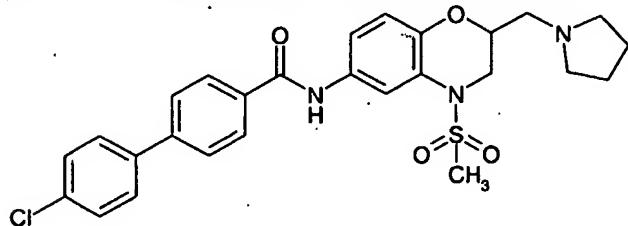
The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 6-amino-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine obtained in Reference Example 117.

5 ¹H-NMR (CDCl₃) δ : 1.60-1.90 (8H, m), 2.50-2.70 (5H, m), 2.71 (2H, d, J=6.3Hz), 2.91-3.00 (2H, m), 3.15-3.21 (1H, brs), 3.42-3.45 (1H, m), 3.77 (1H, brs), 4.15-4.25 (3H, m), 6.20 (1H, s), 6.38 (1H, dd, J=2.1, 8.4 Hz), 6.73 (1H, d, J=8.4 Hz), 6.91 (1H, d, J=2.1 Hz), 6.97-7.03 (2H, m), 7.14-7.19 (2H, m).

10 Melting point: 192-195 °C (crystallization solvent: diisopropyl ether)

Example 251

15 4'-Chloro-N-[4-(methylsulfonyl)-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazin-6-yl][1.1'-biphenyl]-4-carboxamide



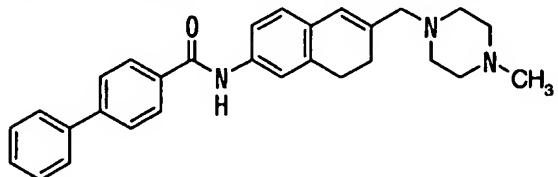
20 The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 6-amino-4-(methylsulfonyl)-2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-1,4-benzoxazine obtained in Reference Example 118.

25 ¹H-NMR (CDCl₃) δ : 1.75-1.85 (4H, m), 2.55-2.70 (4H, m), 2.78 (2H, d, J=6.0Hz), 3.04 (3H, s), 3.27-3.34 (1H, m), 4.24-4.31 (1H, m), 4.31-4.35 (1H, m), 6.98 (1H, d, J=8.7 Hz), 7.45 (2H, d, J=9.0 Hz), 7.50-7.60 (1H, m), 7.53 (2H, d, J=9.0 Hz), 7.67 (2H, d, J=8.4 Hz), 7.84 (1H, s), 7.84 (1H, brs), 7.94 (2H, d, J=8.4 Hz).

30 Melting point: 203-204 °C (crystallization solvent: diisopropyl ether)

Example 252

N-[6-[(4-Methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



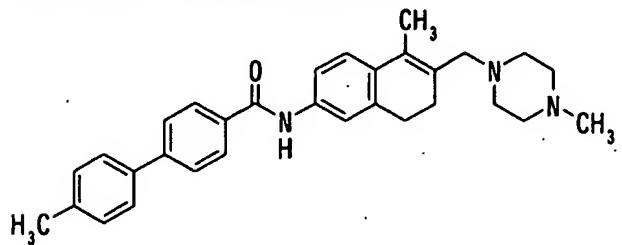
5 The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 106.

10 ^1H NMR (CDCl_3) δ : 2.31 (3H, s), 2.33 (2H, t, $J=8.1$ Hz), 2.49 (8H, bs), 2.84 (2H, t, $J=8.1$ Hz), 3.07 (2H, s), 6.36 (1H, s), 7.02 (1H, d, $J=8.1$ Hz), 7.35-7.52 (5H, m), 7.63 (2H, d, $J=8.1$ Hz), 7.71 (2H, d, $J=8.1$ Hz), 7.80 (1H, s), 7.94 (2H, d, $J=8.1$ Hz).

15 Melting point: 196-198 °C (crystallization solvent:
ethyl acetate)

Example 253

4'-Methyl-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



25 The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

17 ^1H NMR (CDCl_3) δ : 2.08 (3H, s), 2.29 (3H, s), 2.34 (2H, t, $J=7.8$ Hz), 2.42 (3H, s), 2.45 (8H, bs), 2.75 (2H, t, $J=7.8$ Hz), 3.16 (2H, s), 7.26-7.30 (3H, m), 7.44 (1H, d, $J=8.4$

Hz), 7.53-7.55 (3H, m), 7.70 (2H, d, J=8.4 Hz), 8.00 (1H, s), 7.93 (2H, d, J=8.4 Hz).

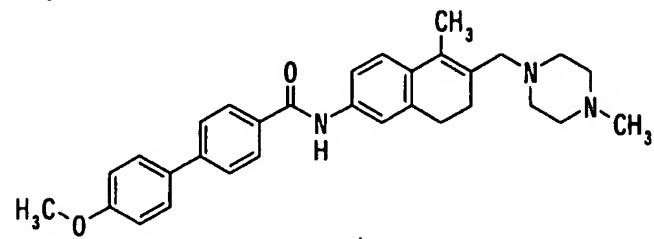
Melting point: 212-214 °C (crystallization solvent: ethyl acetate)

5

Example 254

4'-Methoxy-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

10

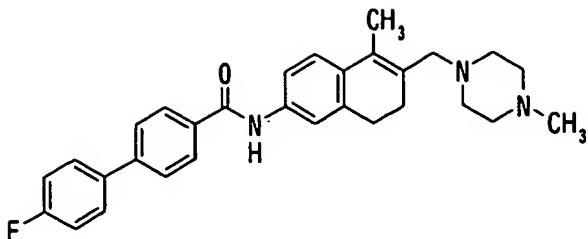


The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

15 ¹H NMR (CDCl₃) δ: 2.08 (3H, s), 2.29 (3H, s), 2.34 (2H, t, J=7.8 Hz), 2.45 (8H, bs), 2.75 (2H, t, J=7.8 Hz), 3.16 (2H, s), 3.87 (3H, s), 7.01 (2H, d, J=8.1 Hz), 7.27 (1H, d, J=8.4 Hz), 7.44 (1H, d, J=8.4 Hz), 7.51 (1H, s), 7.58 (2H, d, J=8.4 Hz), 7.67 (2H, d, J=8.4 Hz), 7.81 (1H, s), 7.92 (2H, d, J=8.4 Hz).

20 Melting point: 215-217 °C (crystallization solvent: ethyl acetate)

25 **Example 255**
4'-Fluoro-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide

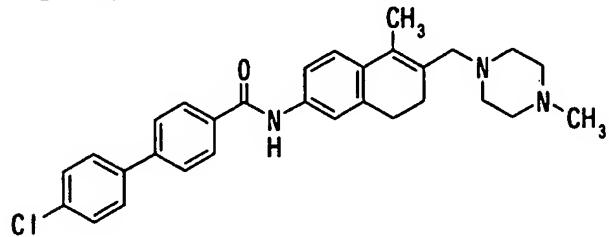


The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

¹H NMR (CDCl₃) δ : 2.08 (3H, s), 2.29 (3H, s), 2.34 (2H, t, J=7.8 Hz), 2.46 (8H, bs), 2.75 (2H, t, J=7.8 Hz), 3.16 (2H, s), 7.17 (2H, d, J=8.4 Hz), 7.28 (1H, d, J=8.4 Hz), 7.44 (1H, d, J=8.4 Hz), 7.51 (1H, s), 7.57-7.62 (2H, m), 7.66 (2H, d, J=8.4 Hz), 7.82 (1H, s), 7.94 (2H, d, J=8.4 Hz).
 Melting point: 233-235 °C (crystallization solvent: ethyl acetate)

Example 256

15 4'-Chloro-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide



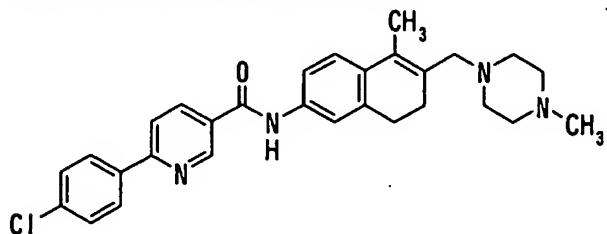
The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

¹H NMR (CDCl₃) δ : 2.08 (3H, s), 2.29 (3H, s), 2.34 (2H, t, J=7.8 Hz), 2.46 (8H, bs), 2.75 (2H, t, J=7.8 Hz), 3.16 (2H, s), 7.28 (1H, d, J=8.4 Hz), 7.43-7.47 (3H, m), 7.51 (1H, s), 7.56 (2H, d, J=8.4 Hz), 7.67 (2H, d, J=8.4 Hz), 7.80 (1H, s), 7.94 (2H, d, J=8.4 Hz).

Melting point: 216-218 °C (crystallization solvent: ethyl acetate)

Example 257

5 6-(4-Chlorophenyl)-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]nicotinamide



The titled compound was obtained by carrying out the
10 same operation as in Reference Example 1, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

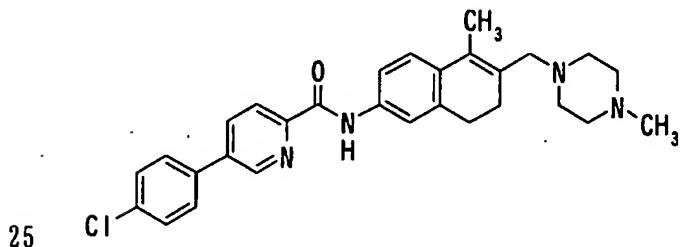
15 ^1H NMR (CDCl_3) δ : 2.09 (3H, s), 2.29 (3H, s), 2.35 (2H, t, $J=8.1$ Hz), 2.46 (8H, bs), 2.75 (2H, t, $J=8.1$ Hz), 3.16 (2H, s), 7.28 (1H, d, $J=8.4$ Hz), 7.43-7.50 (4H, m), 7.83 (2H, d, $J=8.4$ Hz), 8.01 (2H, d, $J=8.4$ Hz), 8.27 (1H, d, $J=8.4$ Hz), 9.13 (1H, s).

Melting point: 219-221 °C (crystallization solvent: ethyl acetate)

20

Example 258

5-(4-Chlorophenyl)-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-2-pyridinecarboxamide



The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 5-

methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

¹H NMR (CDCl₃) δ: 2.09 (3H, s), 2.29 (3H, s), 2.35 (2H, t, J=8.1 Hz), 2.45 (8H, bs), 2.77 (2H, t, J=8.1 Hz), 3.16 (2H, s), 7.30 (1H, d; J=8.1 Hz), 7.49-7.63 (6H, m), 8.05 (1H, dd, J=2.4 Hz, 8.4 Hz), 8.36 (1H, d, J=8.1 Hz), 8.79 (1H, d, J=1.2 Hz), 9.97 (1H, s).

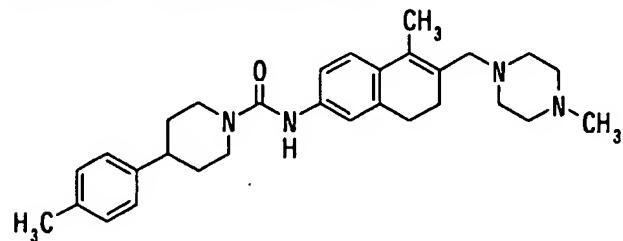
Melting point: 177-179 °C (crystallization solvent: ethyl acetate)

10

Example 259

N-[5-Methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-4-(4-methylphenyl)-1-piperidinecarboxamide

15



15

The titled compound was obtained by carrying out the same operation as in Reference Example 99, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

20

¹H NMR (CDCl₃) δ: 1.60-1.78 (4H, m), 2.05 (3H, s), 2.28 (3H, s), 2.29 (2H, t, J=8.1 Hz), 2.33 (3H, s), 2.46 (8H, bs), 2.65-2.72 (3H, m), 2.93-3.03 (2H, m), 3.13 (2H, s), 4.18-4.23 (2H, m), 6.40 (1H, s), 7.09-7.24 (7H, m).

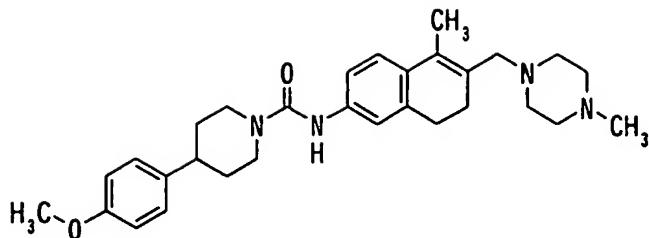
25

Melting point: 176-178 °C (crystallization solvent: ethyl acetate-hexane)

30

Example 260

4-(4-Methoxyphenyl)-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



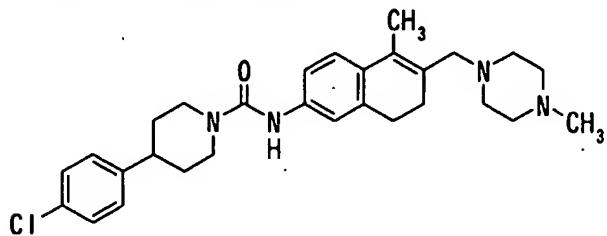
The titled compound was obtained by carrying out the same operation as in Reference Example 99, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

¹H NMR (CDCl₃) δ : 1.68-1.92 (4H, m), 2.05 (3H, s), 2.28 (3H, s), 2.29 (2H, t, J=8.1 Hz), 2.45 (8H, bs), 2.67-2.72 (3H, m), 2.95-3.02 (2H, m), 3.14 (2H, s), 3.80 (3H, s), 4.18-4.22 (2H, m), 6.36 (1H, s), 6.87 (2H, d, J=8.4 Hz), 7.12-7.21 (5H, m).

Melting point: 175-177 °C (crystallization solvent: ethyl acetate)

Example 261

15 4-(4-Chlorophenyl)-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide



The titled compound was obtained by carrying out the same operation as in Reference Example 99, using 5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenamine obtained in Reference Example 115.

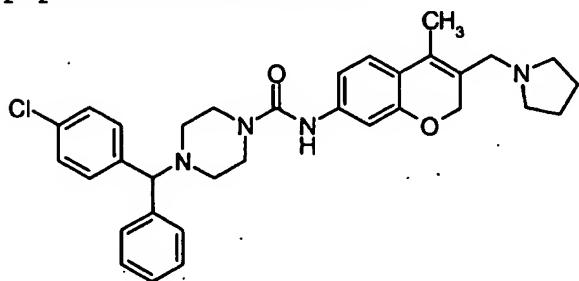
¹H NMR (CDCl₃) δ : 1.67-1.92 (4H, m), 2.05 (3H, s), 2.28 (3H, s), 2.29 (2H, t, J=8.1 Hz), 2.45 (8H, bs), 2.67-2.72 (3H, m), 2.95-3.02 (2H, m), 3.14 (2H, s), 4.18-4.23 (2H, m), 6.36 (1H, s), 7.13-7.30 (7H, m).

Melting point: 141-143 °C (crystallization solvent:

ethyl acetate)

Example 262

5 4-[(4-Chlorophenyl)(phenyl)methyl]-N-[4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]-1-piperazinecarboxamide



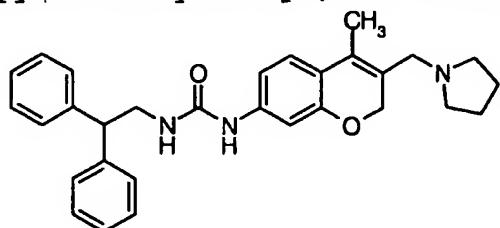
10 The titled compound was obtained by carrying out the same operation as in Reference Example 99, using 4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 107.

15 ^1H NMR (CDCl_3) δ : 1.76 (4H, s), 2.01 (3H, s), 2.42 (4H, t, $J=5.1$ Hz), 2.49 (4H, s), 3.22 (2H, s), 3.48 (4H, t, $J=5.1$ Hz), 4.24 (1H, s), 4.68 (2H, s), 6.23 (1H, s), 6.77 (1H, s), 6.96 (1H, d, $J=8.7$ Hz), 7.09 (1H, d, $J=8.7$ Hz), 7.19-7.61 (9H, m).

Melting point: 104-106 °C (crystallization solvent: ethyl acetate - n-hexane)

20 **Example 263**

N-(2,2-Diphenylethyl)-N'-(4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl)urea



25 The titled compound was obtained by carrying out the same operation as in Reference Example 99, using 4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine

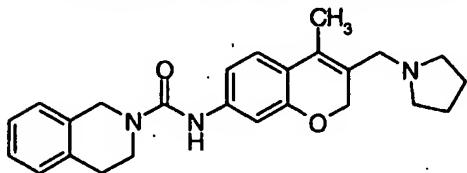
obtained in Reference Example 107.

¹H NMR (CDCl₃) δ: 1.76 (4H, s), 1.99 (3H, s), 2.49 (4H, s), 3.22 (2H, s), 3.83 (2H, t, J=7.8 Hz), 4.18 (1H, t, J=7.8 Hz), 4.66 (2H, s), 4.96 (1H, s), 6.48 (1H, s), 6.57 (1H, s), 6.69 (1H, d, J=8.1 Hz), 6.98 (1H, d, J=8.1 Hz), 7.20-7.30 (10H, m).

Melting point: 166-168 °C (crystallization solvent: ethyl acetate - n-hexane)

10 Example 264

N-[4-Methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]-3,4-dihydro-2(1H)-isoquinolinecarboxamide



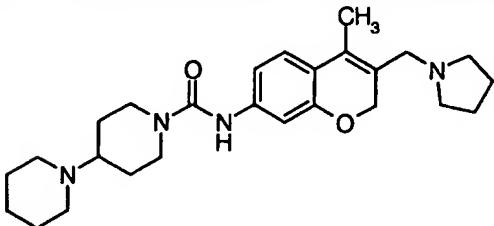
15 The titled compound was obtained by carrying out the same operation as in Reference Example 99, using 4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 107.

¹H NMR (CDCl₃) δ: 1.76 (4H, s), 2.02 (3H, s), 2.49 (4H, s), 2.92 (2H, t, J=6.0 Hz), 3.23 (2H, s), 3.71 (2H, t, J=6.0 Hz), 20 4.65 (2H, s), 4.68 (2H, s), 6.43 (1H, s), 6.86 (1H, d, J=1.8 Hz), 7.02-7.22 (6H, m).

Melting point: 135-137 °C (crystallization solvent: ethyl acetate - n-hexane)

25 Example 265

N-[4-Methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]-4-(1-piperidinyl)-1-piperidinecarboxamide



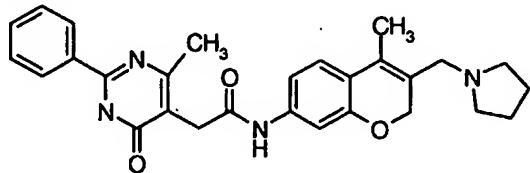
The titled compound was obtained by carrying out the same operation as in Reference Example 99, using 4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 107.

5 ¹H NMR (CDCl₃) δ : 1.27-1.89 (14H, m), 2.02 (3H, s), 2.49-2.51 (9H, m), 2.83-2.90 (2H, m), 3.23 (2H, s), 4.08-4.12 (2H, m), 4.68 (2H, s), 6.31 (1H, s), 6.80 (1H, d, J=2.4 Hz), 6.98 (1H, dd, J=2.4 Hz, 8.4 Hz), 7.09 (1H, d, J=8.4 Hz).

10 Melting point: 98-100 °C (crystallization solvent: ethyl acetate - n-hexane)

Example 266

15 2-(4-Methyl-6-oxo-2-phenyl-1,6-dihydro-5-pyrimidinyl)-N-[4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]acetamide



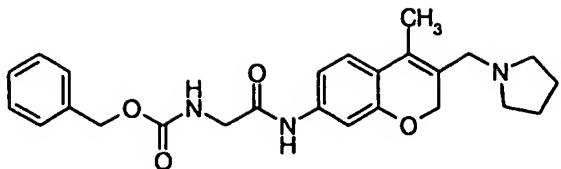
20 The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 107.

25 ¹H NMR (CDCl₃) δ : 1.76 (4H, s), 1.98 (3H, s), 2.49 (4H, s), 2.61 (3H, s), 3.22 (2H, s), 3.65 (2H, s), 4.65 (2H, s), 6.86-7.00 (4H, m), 7.54 (3H, s), 8.01 (2H, s), 8.87 (1H, s).

25 Melting point: 255-257 °C (crystallization solvent: ethyl acetate - n-hexane)

Example 267

30 Benzyl 2-[[4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-yl]amino]-2-oxoethylcarbamate



The titled compound was obtained by carrying out the same operation as in Reference Example 1, using 4-methyl-3-(1-pyrrolidinylmethyl)-2H-chromen-7-amine obtained in Reference Example 107.

¹H NMR (CDCl₃) δ : 1.78 (4H, s), 2.03 (3H, s), 2.53 (4H, s), 3.26 (2H, s), 3.99 (2H, d, J=4.8 Hz), 4.71 (2H, s), 5.17 (2H, s), 5.50 (1H, bs), 7.00-7.14 (4H, m), 7.36 (5H, s), 7.80 (1H, bs).

Melting point: 143-145 °C (crystallization solvent: ethyl acetate - n-hexane)

Preparation Example 1

| | |
|---|--------------|
| (1) Compound obtained in | |
| Reference Example 25 | 50 mg |
| (2) Lactose | 34 mg |
| (3) Corn starch | 10.6 mg |
| (4) Corn starch (paste) | 5 mg |
| (5) Magnesium stearate | 0.4 mg |
| <u>(6) Carboxymethylcellulose calcium</u> | <u>20 mg</u> |
| | Total 120 mg |

In accordance with a conventional manner, the above (1) to (6) are admixed and tableted using a tabletting machine to give tablets.

25

Preparation Example 2

| | |
|---|--------------|
| (1) Compound obtained in Example 1 | 50 mg |
| (2) Lactose | 34 mg |
| (3) Corn starch | 10.6 mg |
| (4) Corn starch (paste) | 5 mg |
| (5) Magnesium stearate | 0.4 mg |
| <u>(6) Carboxymethylcellulose calcium</u> | <u>20 mg</u> |
| | Total 120 mg |

In accordance with a conventional manner, the above (1) to (6) are admixed and tableted using a tabletting machine to give tablets.

5 Reference Example 1-1

Amplification of rat SLC-1 receptor cDNA by PCR method using rat-brain-originated cDNA

Reverse transcription reaction was done using random primer, with rat-brain-originated poly (A)'RNA (Clone Tech Co.) used as a template. Reagent from the TaKaRa RNA PCR ver. 2 kit was used for the reverse transcription reaction.

Next, using this reverse transcription product as a template, amplification was done by a PCR method using synthetic DNA primers with sequence numbers 1 and 2.

15 Synthetic DNA primer was constructed to amplify genes in the domain where genes are translated by receptor protein.

At that time, individual restriction enzyme recognition sequences were also added on the 5' side and 3' side of the gene, to add a nucleotide sequence on the 5' side of gene which recognized restriction enzyme Sal I, and to add a nucleotide sequence on the 3' side of the gene which recognized the restriction enzyme Spe I. The reactant was constituted of 5 µl of cDNA template, 0.4 µM of synthetic DNA primer, 0.25 mM of dNTPs, 0.5 µl of Pfu (StrataGene Co.) DNA polymerase, and buffers attached to enzymes, with total reaction quantity set at 50 µl.

30 A thermal cycler (Parkin Elmer Co.) was used to produce cycles for amplification. After heating at 94°C for 60 seconds, the cycle consisting of 94°C for 60 seconds, 60°C for 30 seconds, and 72°C for 150 seconds, was repeated 35 times, and finally reaction was conducted at 72°C for 10 minutes. After 0.8% agarose gel electrophoresis, the amplified products were confirmed by ethidium bromide dying.

35

Reference Example 1-2

Subcloning of PCR products into plasmid vector, and confirmation of an amplified cDNA sequence by decoding of a nucleotide sequence in an inserted cDNA portion

The reaction product after PCR conducted in Reference 5 Example 1-1 was separated using 0.8% low-melting point agarose gel. After the band section was cut out using a razor, DNA was recovered by conducting fragmentation, phenol extraction, phenol-chloroform extraction and ethanol precipitation. The recovered DNA was subcloned on 10 plasmid vector PCR-Script Amp SK(+) in accordance with prescription of the PCR-Script™ Amp SK(+) cloning kit (Stratagene Co.). After this was introduced into Escherichia coli XL-1 Blue (Stratagene Co.) by transformation, the clones with fragments of inserted cDNA 15 were selected in LB agar culture medium containing ampicillin and X-gal. Only clones showing white color were separated using a sterilized toothpick, and transformant E. coli XL-1 Blue/rat SLC-1 was obtained.

Each clone was cultured overnight in LB culture medium 20 containing ampicillin, and plasmid DNA was prepared using QIA prep8 mini prep (Qiagen). A portion of the prepared DNA was digested with Sal I and Spe I, and the size of the inserted receptor cDNA fragment was confirmed. Reactions to determine nucleotide sequences were carried out using 25 a DyeDeoxy Terminator Cycle Sequence Kit (Parkin Elmer Co.), and decoded using a fluorescent light automatic sequencer. The sequences of the 3 clones obtained were analyzed, and it was confirmed that all of them match the reported gene sequence (Sequence number: 4) in which the 30 Sal I recognition sequence is added on the 5' side and the Spe I recognition sequence is added on the 3' side of the cDNA sequence (Lakaye, B., et al., Biochim. Biophys. Acta, Vol. 1401, pp. 216-220 (1998), accession No. AF08650) coding rat SLC-1 protein (Sequence number: 3).

Preparation of CHO cells for rat SLC-1 expression

The full-length amino acid sequence of rat brain originated SLC-1, which was confirmed in Reference Example 1-2, was coded, and plasmid was prepared using a plasmid 5 Midi Kit (Qiagen) from the E. coli transformed by the plasmid, to which the gene with Sal I recognition sequence added to the 5' side and Spe I recognition sequence added to the 3' side, had been introduced. Then, the insert section was cut out by digesting with Sal I and Spe I. The 10 insert DNA was cut out with a razor from the agarose gel after electrophoresis.

Next, fragmentation, phenol extraction, phenol-chloroform extraction, and ethanol precipitation, were conducted and the DNA was recovered. This insert DNA was 15 added to vector plasmid pAKKO-111H (the same vector plasmid as pAKKO1.11H described in Hinuma, S., et al., Biochim. Biophys. Acta, Vol. 1219, pp. 251-259 (1994)) for animal cell expression which was digested with Sal I and Spe I, and ligation was conducted using T4 ligase (TaKaRa Shuzo), 20 to construct pAKKO-SLC-1 plasmid for protein expression.

After E. coli DH5 transformed by pAKKO-SLC-1 was cultured, pAKKO-SLC-1 plasmid DNA was prepared using a Plasmid Midi Kit (Qiagen). This was introduced into CHO dhfr⁻ cells in accordance with the attached protocol, using 25 a CellPfect Transfection Kit (Amersham Pharmacia Biotech Co.). A coprecipitating suspension of 10 µg of DNA and calcium phosphate was prepared, and this suspension was added to 10 cm Petri dishes in which 5 × 10⁵ or 1 × 10⁶ of CHO dhfr⁻ cells had been seeded 24 hours previously. After 30 these cells were cultured for 1 day in MEMa culture medium containing 10% fetal bovine serum, subculture was conducted, and cultivation was conducted in selective culture medium, MEMa culture medium containing no nucleic acid but containing 10% dialyzed fetal bovine serum. 56 35 clones of colonies of the transformed CHO cells expressing SLC-1, proliferated in the selective culture medium, were

selected.

Reference Example 1-4

Selection of CHO/SLC-1 cell strain expressing a large
5 quantity of full-length rat SLC-1 receptor protein mRNA
The quantity of expressed full-length rat SLC-1
receptor protein mRNA of 56 clones of the CHO/SLC-1 strains
established in Reference Example 1-3, was measured using
a Cytostar T Plate (Amersham Pharmacia Biotech Co.) as shown
10 below according to the attached protocol. Each well of the
Cytostar T Plate was seeded with each clone of the CHO/SLC-1
strain by 2.5×10^4 , and cultured for 24 hours, then the
cells were fixed using 10% formalin. After 0.25% Triton
X-100 was added to each well to increase cell permeability.
15 35 S-labeled riboprobes with sequence number: 5 were added
and hybridized. 20 mg/ml of RNaseA was added to each well
to digest free riboprobes. After the plate was thoroughly
washed, the radioactivity of the hybridized riboprobes was
determined using a Topcounter. Strains with high
20 radioactivity showed large amounts of mRNA expression. In
particular, mainly used was Clone number 44 among 3 clones
which showed large amounts of mRNA expression.

Reference Example 1-5

25 Isolation of plasmid containing human SLC-1 cDNA

After nicks were inserted into the DNA of Human fetal
brain originated cDNA library (SUPERSCRIPT™ cDNA Library;
GIBCOBRL Co.) according to the manual of the Genetranner
cDNA positive selection system (GIBCOBRL Co.), using phage
30 F1 endonuclease, single stranded human fetal brain
originated cDNA library was prepared by digesting the
above-mentioned library with Escherichia coli exonuclease
III.

Biotin-14-dCTP was added to the 3' end of synthetic
35 oligonucleotide (equivalent to 1434-1451 of accession No.
U71092), sequence number: 6 which was prepared according

to the report by Kolakowski Jr., et al. (Kolakowski Jr., et al. (1996) FEBS Lett. Vol. 398, pp. 253-258) using Terminal Deoxynucleotidyl Transferase, and biotinylated oligonucleotide was prepared. The above manual was
5 followed regarding composition of a reaction mixture and reaction time.

After 4 µg of single stranded human fetal brain originated cDNA library was kept at 95°C for 1 minute, the library was rapidly cooled on ice. 20 ng of biotinylated
10 oligonucleotide was added, which was hybridized using the attached hybridization buffer at 37°C for 1 hour. Streptoavidin beads were added to the mixture, then single stranded human fetal brain originated cDNA hybridized by biotinylated oligonucleotide, was isolated using a MAGNA-
15 SEP Magnetic Particle Separator (GIBCOBRL Co.). The complementary strand was synthesized according to the manual, using as primer 50 ng of synthetic oligonucleotide (equivalent to 1011 - 1028 of accession No. U71092) of sequence number: 7, prepared based on the report by
20 Kolakowski Jr., et al (Kolakowski Jr., et al. (1996) FEBS Lett. Vol. 398, pp. 253-258), to give the double stranded plasmid.

Reference Example 1-6

25 Determination of nucleotide sequence of plasmid containing isolated human SLC-1 cDNA

After the plasmid obtained in Reference Example 1-5 was introduced into ELECTROMAX™DH10B™ Cells by the electroporation method, clones with cDNA inserted
30 fragments were selected in LB agar culture medium containing ampicillin and X-gal. Using a sterilized toothpick, only the clones showing white color were separated to give transformant E. coli DH10B/hSLC-1. Individual clones were cultured overnight in LB culture
35 medium containing ampicillin, and the plasmid DNA was refined using QIA prep8 mini prep (Qiagen). The reactions

to determine nucleotide sequence were conducted using a DyeDeoxy Terminator Cycle Sequence Kit (Parkin Elmer Co.), and the nucleotide sequence was decoded using a fluorescent light automatic sequencer.

5 As the results, obtained was the sequence shown in Sequence number: 8. The amino acid sequence (Sequence number: 9) coded by the nucleotide sequence obtained here, differs from the human SLC-1 amino acid sequence predicted as the sequence analogized from rat SLC-1 based on human
10 chromosome DNA sequence (accession number: Z86090) containing human SLC-1 sequence, in the report by Lakaye, et al. (Lakaye, B., et al. (1998) Biochim. Biophys. Acta. Vol. 1401, pp. 216-220). This shows the presence of ATG, the initiation codon, on mRNA, in the 69 and 64 amino acids
15 upstream from the estimated sequence. Escherichia coli DH10B/phSLC1L8, the transformant produced by the plasmid containing DNA coding this sequence was deposited at IFO and NIBH.

20 Reference Example 1-7

Amplification of human SLC-1cDNA by PCR method using human fetal brain originated cDNA

Amplification by the PCR method was conducted using as the template plasmid containing human SLC-1 DNA sequence
25 cloned by the gene trap method, and using synthetic DNA primers of sequence number: 10 and sequence number: 11, and synthetic DNA primers of sequence number: 12 and sequence number: 13, respectively. The former amplified DNA and the latter amplified DNA were named as "human SLC-1(S)" and
30 "human SLC-1(L)", respectively. The synthetic DNA primer was constructed so that the genes in the domain translated to the receptor protein were amplified. At that time, a recognition sequence for each restriction enzyme was added on the 5' side and 3' side, so that the nucleotide sequence
35 recognized by restriction enzyme Sal I would be added on the 5' side of the gene, and the nucleotide sequence

recognized by restriction enzyme Spe I would be added on the 3' side. The composition of the reaction mixture for human SLC-1(S) amplification was: 5 µl of plasmid template containing human SLC-1 DNA sequence, 0.4 µM of respective synthetic DNA primers, 0.2 mM of dNTPs and 0.5 µl of Pfu DNA polymerase and buffers attached to the enzyme, with total quantity for reaction set at 50 µl. A thermal cycler (Parkin Elmer Co.) was used for the cycles for amplification. After heating at 94°C for 60 seconds, the cycle consisting of 94°C for 60 seconds, 57°C for 60 seconds, and 72°C for 150 seconds, was repeated 25 times, and finally the temperature of the reactant was maintained at 72°C for 10 minutes. The composition of the reaction mixture for human SLC-1(L) amplification was 5 µl of plasmid template containing human SLC-1 DNA sequence, 0.4 µM of respective synthetic DNA primers, 0.2 mM of dNTPs, 0.5 µl of Pfu DNA polymerase and buffers attached to the enzymes, with total quantity for reaction set at 50 µl. A thermal cycler (Parkin Elmer Co.) was used for the cycles for amplification. After heating at 94°C for 60 seconds, the cycle consisting of 94°C for 60 seconds, 60°C for 60 seconds, and 72°C for 3 minutes, was repeated 25 times, and finally the temperature of the reactant was maintained at 72°C for 10 minutes. After 0.8% agarose gel electrophoresis, confirmation of amplified products was conducted by ethidium bromide dying.

Reference Example 1-8

Subcloning of PCR product into plasmid vector and confirmation of amplified cDNA sequence by decoding of nucleotide sequence of inserted cDNA section

The reaction product after PCR in Reference Example 1-7 was separated using 0.8% low-melting point agarose gel, and the band section was cut out using a razor. After that, fragmentation, phenol extraction, phenol-chloroform extraction, and ethanol precipitation were conducted, and

the DNA was recovered. The recovered DNA was subcloned into pCR-Script Amp SK(+) plasmid vector, as prescribed by the PCR-Script™ Amp SK(+) cloning kit (Stratagene Co.). After this was introduced into Escherichia coli DH5a competent 5 cells (TOYOBO) and transformed, the clones with cDNA inserted fragments were selected in LB agar culture medium containing ampicillin and X-gal. Using a sterilized toothpick, only clones showing white color were separated to give E. coli DH5a/hSLC-1(S), which is a transformant of 10 human SLC-1 (S), and E. coli DH5a/hSLC-1(L), which is a transformant of human SLC-1 (L). Each clone was cultured overnight in LB culture medium containing ampicillin, and plasmid DNA was prepared using QIA prep8 mini prep (Qiagen). Some of the prepared DNA was digested with Sal I and Spe 15 I restriction enzymes, and the size of the receptor cDNA fragments inserted was confirmed. The reactions to determine nucleotide sequence were conducted using a DyeDeoxy Terminator Cycle Sequence Kit (Parkin Elmer Co.) and the nucleotide sequence was decoded using a fluorescent 20 light automatic sequencer. The sequence of the obtained clones respectively matched the DNA sequence (sequence number:14) which should be amplified by synthetic DNA primers of sequence number: 10 and sequence number: 11 using human SLC-1 gene as a template, and the DNA sequence 25 (sequence number: 15) which should be amplified by synthetic DNA primers of sequence number: 12 and sequence number: 13 using human SLC-1 gene as a template.

Reference Example 1-9

30 Preparation of CHO cells for expression of human SLC-1(S), and CHO cells for expression of human SLC-1(L)

Plasmid was prepared from the E. coli clones transformed by the plasmid wherein inserted were human SLC-1(S) and human SLC-1(L) whose sequences were confirmed 35 in Reference Example 1-8, using a Plasmid Midi Kit (Qiagen), and the insert section was cut out using Sal I and Spe I

restriction enzymes. After electrophoresis was conducted, the insert DNA was cut out from agarose gel using a razor. Next, fragmentation, phenol extraction, phenol-chloroform extraction, and ethanol precipitation 5 were conducted, and the insert DNA was recovered.

This insert DNA was added to pAKKO-111H vector plasmid for animal cell expression, digested with Sal I and Spe I (the same vector plasmid as the pAKKO1.11H described in Hinuma, S., et al., Biochim. Biophys. Acta, Vol. 1219, pp. 10 251-259 (1994)), and ligation was conducted by adding T4 ligase (TaKaRa Shuzo), to construct pAKKO-hSLC-1(S) and pAKKO-hSLC-1(L) plasmids for protein expression.

After E. coli DH5 α (TOYOBO) transformed by pAKKO-hSLC-1(S) and pAKKO-hSLC-1(L) was cultured, pAKKO-hSLC-15 1(S) and pAKKO-hSLC-1(L) plasmid DNAs were prepared using a Plasmid Midi Kit (Qiagen). These were introduced into CHO dhfr $^{-}$ cells in accordance with the attached protocol, using a CellPfect Transfection Kit (Amersham Pharmacia Biotech Co.). A coprecipitative suspension of 10 μ g of DNA 20 with calcium phosphate was made, which was added to 10 cm Petri dishes seeded 24 hours in advance with 5×10^5 or 1×10^6 CHO dhfr $^{-}$ cells. After the above was cultured for 1 day in MEM α culture medium containing 10% fetal bovine serum, subculture was conducted, and then cultivation was 25 conducted in MEM α culture medium containing no nucleic acid but containing 10% dialyzed fetal bovine serum, which is a selective culture medium. 56 clones of colonies of transformed cells which are human SLC-1(S) gene introduced CHO cells, and 61 clones of colonies of transformed cells 30 which are human SLC-1(L) gene introduced CHO cells, both of which proliferated in the selective culture medium, were selected.

Reference Example 1-10

35 Selection of cell colonies into which genes with large quantities of human SLC-1(S) and human SLC-1 (L) mRNA

expression have been introduced

The quantities of expressed mRNA of 56 clones of CHO/hSLC-1(S) colonies and 61 clones of CHO/hSLC-1(L) colonies, both of which were established in Reference 5 Example 1-9, were measured in accordance with the attached protocol using a Cytostar T Plate (Amersham Pharmacia Biotech Co.) as shown below.

After each well of the Cytostar T Plate was seeded with each clone of CHO/hSLC-1(S) colonies and CHO/hSLC-1(L) 10 colonies by 2.5×10^4 , and cultured for 24 hours, the cells were fixed using 10% formalin.

After 0.25% Triton X-100 was added to each well to increase cell permeability, 35 S-labeled riboprobe of sequence number: 16 was added and hybridization was 15 conducted.

20 mg/ml of RNaseA was added to each well to digest free riboprobe. After the plate was washed well, the radioactivity of the hybridized riboprobe was determined. Colonies showing high radioactivity expressed large 20 quantities of mRNA. Of the 7 clones which expressed large quantities of mRNA, mainly used was Clone number 57.

Experimental Example 1

Determination of antagonist activity using GTPgS binding 25 assay of test compound

Membrane fraction was prepared by the following method, using the human SLC-1 expressing CHO cell clone 57 obtained in Reference Example 1-10, and the rat SLC-1 expressing CHO cell clone 44 obtained in Reference Example 30 1-4.

The human and rat SLC-1 expressing CHO cells (1×10^8) were scraped in buffer saline phosphate (pH 7.4) to which 5 mM EDTA (ethylenediaminetetraacetic acid) had been added, and centrifuged. 10 ml of homogenized buffer (10 mM NaHCO₃, 35 5 mM EDTA, pH 7.5) was added to the cell pellets, and they were homogenized using a Polytron homogenizer. The

supernatant obtained by centrifugation at $400 \times g$ for 15 minutes was further centrifuged at $100,000 \times g$ for 1 hour, to obtain the membrane fraction precipitate. This precipitate was suspended in 2 ml of assay buffer [50 mM Tris-HCl(pH 7.5), 1 mM EDTA, 0.1% BSA (bovine serum albumin), 10 mM MgCl₂, 100 mM NaCl, 1 μ M GDP (guanosine 5'-diphosphate), 0.25 mM PMSF (phenylmethylsulfonyl fluoride), 1 mg/ml pepstatin, 20 mg/ml leupeptin, 10 mg/ml phosphoramidon], which was centrifuged at $100,000 \times g$ for 1 hour. The membrane fraction recovered as precipitate was suspended again in 2 ml of assay buffer, and after the suspension was divided, individual portions were preserved at -80°C and thawed before every use.

Determination of antagonist activity of the test compound was conducted as shown below. After 171 μ l of SLC-1 expressing CHO cell membrane fractions diluted with assay buffer was poured into each well of a 96-well polypropylene plate, 2 μ l of 3×10^{-10} M MCH diluted with DMSO solution, 2 μ l of test compound solution diluted to various concentrations, and 25 μ l of [³⁵S]-Guanosine 5'-(γ -thio) triphosphate (produced by Daiichi Kagaku Yakuhin) were added respectively. (Final concentration of cell membrane: 20 μ g/ml, final concentration of [³⁵S]-Guanosine 5'-(γ -thio) triphosphate: 0.33 nM).

After this reaction mixture was allowed to react for 1 hour under stirring, it was filtered under vacuum using a glass filter (GF-C), then the filter was washed 3 times with 300 μ l of washing solution (50 mM Tris-HCl buffer solution pH 7.5). 50 ml of liquid scintillator was added to the glass filter, and residual radioactivity was determined using a liquid scintillation counter.

The IC₅₀ value of the compound was calculated from the binding inhibition rate (%), based on the definition that the binding inhibition rate (%) = (radioactivity when compound and MCH were added - radioactivity when DMSO solution was added)/(radioactivity when MCH was added -

radioactivity when DMSO solution was added) × 100.

The results were shown below.

| Compound Number | Inhibition Activity (IC ₅₀ value: nM) |
|----------------------|---|
| Reference Example 25 | 90 |
| Example 1 | 40 |

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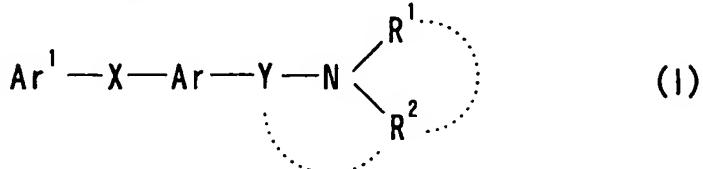
Industrial Applicability

Compounds (I), (I') and salts thereof possess excellent MCH receptor antagonistic activities, and are useful as an agent for preventing or treating obesity, etc.

10

CLAIMS

1. A melanin-concentrating hormone antagonist which comprises a compound of the formula :



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wherein Ar¹ is a cyclic group which may have substituents; X is a spacer having a main chain of 1 to 6 atoms; Y is a bond or a spacer having a main chain of 1 to 6 atoms; Ar is a monocyclic aromatic ring which may be condensed with 10 a 4 to 8 membered non-aromatic ring, and may have further substituents; R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing 15 hetero ring which may have substituents; R² may form a spiro ring together with Ar; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof.

20 2. An antagonist according to claim 1, wherein Y is a spacer having a main chain of 1 to 6 atoms; R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring 25 which may have substituents; or R² may form a spiro ring together with Ar.

30 3. An antagonist according to claim 2, wherein Ar¹ is an aromatic group which may have substituents; and "a hydrocarbon group which may have substituents" for R¹ and R² is "C₁₋₆ alkyl which may have substituents".

4. An antagonist according to claim 1, wherein the cyclic

group for Ar¹ is C₆₋₁₄ monocyclic or condensed polycyclic aromatic hydrocarbon group.

5. An antagonist according to claim 1, wherein the cyclic group for Ar¹ is a group formed by removing an optional one hydrogen atom from an aromatic ring assemble in which 2 or 3 C₆₋₁₄ monocyclic or condensed polycyclic aromatic hydrocarbon groups are directly bonded by single bonds.

10 6. An antagonist according to claim 1, wherein the cyclic group for Ar¹ is a group formed by removing an optional one hydrogen atom from an aromatic ring assemble in which C₆₋₁₄ monocyclic or condensed polycyclic aromatic hydrocarbon and 5 to 10 membered aromatic hetero ring are directly bonded by a single bond.

20 7. An antagonist according to claim 1, wherein Ar¹ is phenyl, biphenylyl, phenyl-pyridyl, phenyl-furyl, phenyl-isoxazolyl, diphenyl-oxazolyl, pyridyl-phenyl, phenyl-pyrimidinyl, benzofuranyl-phenyl, furyl-phenyl, terphenyl, thienyl-phenyl, indolyl, naphthyl-oxadiazolyl, benzofuranyl-oxadiazolyl, benzothienyl, benzofuranyl, fluorenyl, pyridyl-pyrrolyl or thioxanthenyl;

25 each of which may have 1 to 3 substituents selected from the group consisting of halogen atom; nitro; C₁₋₃ alkylenedioxy; optionally halogenated C₁₋₆ alkyl; hydroxy-C₁₋₆ alkyl; optionally halogenated C₃₋₆ cycloalkyl; optionally halogenated C₁₋₆ alkoxy; optionally halogenated C₁₋₆ alkythio; hydroxy; C₇₋₁₉ aralkyloxy which may have substituents; C₆₋₁₄ aryloxy which may have substituents; amino; mono-C₁₋₆ alkylamino; di-C₁₋₆ alkylamino; 5 to 7 membered saturated cyclic amino which may have substituents and may be condensed with a benzene ring; 5 to 7 membered 30 non-aromatic heterocyclic groups which may have substituents; formyl; carboxy; C₆₋₁₄ aryl-carbonyl which may

35

have substituents; C₆₋₁₄ aryl-carbamoyl which may have substituents; aromatic hetero ring-carbamoyl which may have substituents; C₁₋₆ alkoxy-carbonyl; optionally halogenated C₁₋₆ alkyl-carboxamide; C₆₋₁₄ aryl-carboxamide which may have substituents; C₇₋₁₉ aralkyl-carboxamide which may have substituents; aromatic hetero ring-carboxamide which may have substituents; N-(C₆₋₁₄ aryl-carbonyl which may have substituents)-N-C₁₋₆ alkylamino; C₆₋₁₄ arylamino-carbonylamino which may have substituents; C₆₋₁₄ 10 arylsulfonylamino which may have substituents; C₆₋₁₄ aryl-carbonyloxy which may have substituents; oxo; carboxy-C₁₋₆ alkyl; C₁₋₆ alkoxy-carbonyl-C₁₋₆ alkyl; C₇₋₁₉ aralkyl which may have substituents; aromatic hetero ring-C₁₋₆ alkoxy; and cyano.

15

8. An antagonist according to claim 1, wherein Ar¹ is piperidinyl, piperazinyl, pyrrolidinyl, dihydropyridyl or tetrahydropyridyl; each of which may have 1 or 2 substituents selected from the group consisting of oxo, C₆₋₁₄ 20 aryl which may have substituents, hydroxy, C₇₋₁₉, aralkyloxy-carbonyl, and C₇₋₁₉ aralkyl.

9. An antagonist according to claim 1, wherein the "spacer having a main chain of 1 to 6 atoms" for X and Y 25 is a bivalent group consisting of 1 to 3 species selected from -O-, -S-, -CO-, -SO-, -SO₂-, -NR⁸- (R⁸ is hydrogen atom, optionally halogenated C₁₋₆ alkyl, optionally halogenated C₁₋₆ alkyl-carbonyl, optionally halogenated C₁₋₆ 30 alkylsulfonyl), and a bivalent C₁₋₆ non-cyclic hydrocarbon group which may have substituents.

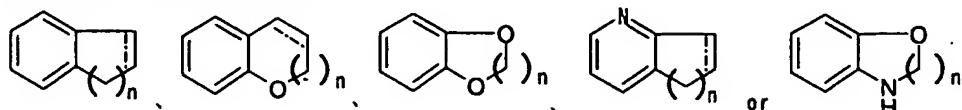
10. An antagonist according to claim 1, wherein X is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}- 35 wherein R^{8c} is hydrogen atom or C₁₋₆ alkyl.

35

11. An antagonist according to claim 1, wherein Y is an

optionally halogenated bivalent C₁₋₆ non-cyclic hydrocarbon group.

12. An antagonist according to claim 1, wherein Ar is a
5 ring of the formula :



wherein ----- is a single bond or double bond, n is an integer of 1 to 4.

10 13. An antagonist according to claim 1, wherein R¹ and R² are hydrogen atom or C₁₋₆ alkyl which may have substituents; or R¹ and R², together with the adjacent nitrogen atom, form a 3 to 8 membered nitrogen-containing hetero ring.

15 14. An antagonist according to claim 1, which is an agent for preventing or treating diseases caused by a melanin-concentrating hormone.

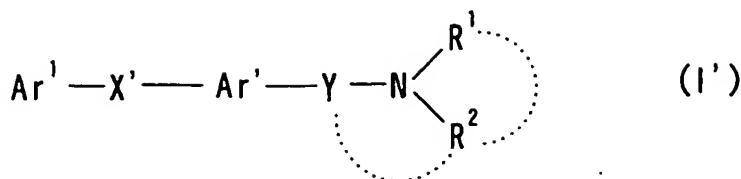
20 15. An antagonist according to claim 1, which is an agent for preventing or treating obesity.

16. An antagonist according to claim 1, which is an anorectic agent.

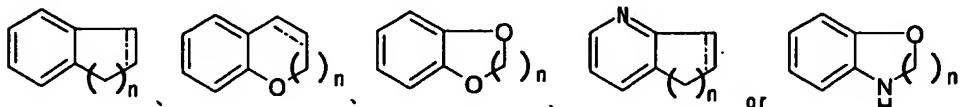
25 17. A pharmaceutical, which comprises a melanin-concentrating hormone antagonist in combination with at least one species selected from the group consisting of an agent for treating diabetes, an agent for treating hypertension and an agent for treating arteriosclerosis.

30

18. A compound of the formula :

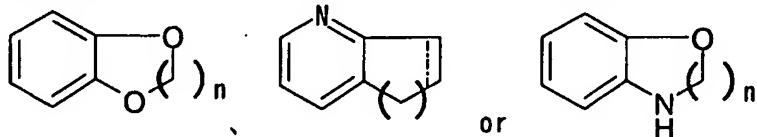


wherein Ar¹ is a cyclic group which may have substituents;
Ar' is a ring of the formula :



- 5 wherein ----- is a single bond or double bond, n is an integer of 1 to 4, and each ring may have substituents; X' is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl;
- Y is a spacer having a main chain of 1 to 6 atoms;
- 10 R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R², together with the adjacent nitrogen atom and Y, may form a
- 15 nitrogen-containing hetero ring which may have substituents;

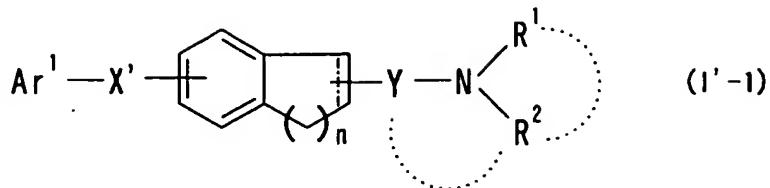
provided that Ar' is a ring of the formula :



wherein symbols have the same meanings as defined above,

- 20 and each ring may have substituents, when X' is -SO₂NH-; ... and provided that Ar¹ is not biphenylyl which may be substituted, when X' is -CONH- and Ar' is any one of benzopyran, dihydrobenzopyran, dihyrobenzoxazine, dihydrobenzoxazole or tetrahydrobenzoxazepine;
- 25 (excluding N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-4-biphenylylcarboxamide); or a salt thereof.

19. A compound of the formula :



wherein Ar^1 is a cyclic group which may have substituents;

--- is a single bond or double bond;

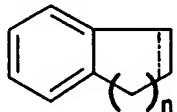
n is an integer of 1 to 4;

5 X' is $-\text{CONR}^{8c}-$, $-\text{NR}^{8c}\text{CO}-$ or $-\text{CH}=\text{CH}-\text{CONR}^{8c}-$ where R^{8c} is hydrogen atom or C_{1-6} alkyl;

Y is a spacer having a main chain of 1 to 6 atoms;

R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 , together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R^2 , together with the adjacent nitrogen atom and Y , may form a nitrogen-containing hetero ring which may have substituents;

15 a ring of the formula :

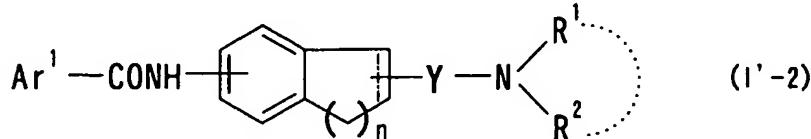


wherein symbols have the same meanings as defined above, may have further substituents;

provided that $\text{N}-(2-(\text{N},\text{N}-\text{dimethylamino})\text{methyl}-6-$

20 tetralinyl)-4-biphenylcarboxamide is excluded; or a salt thereof.

20. A compound according to claim 19, which is of the formula :



25

wherein R^1 and R^2 are independently hydrogen atom or a hydrocarbon group which may have substituents; R^1 and R^2 ,

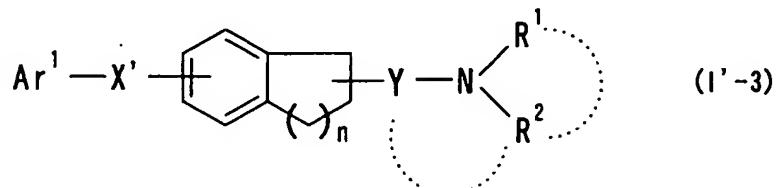
together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; the other symbols have the same meanings as defined in claim 19.

5

21. A compound according to claim 20, wherein Ar¹ is an aromatic group which may have substituents; and "a hydrocarbon group which may have substituents" for R¹ and R² is "C₁₋₆ alkyl which may have substituents".

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22. A compound of the formula :

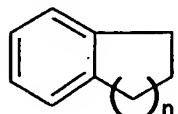


wherein Ar¹ is a cyclic group which may have substituents; n is an integer of 1 to 4;

15 X' is -CONR^{8c}-, -NR^{8c}CO- or -CH=CH-CNR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl;

Y is a spacer having a main chain of 1 to 6 atoms; R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;

25 a ring of the formula :

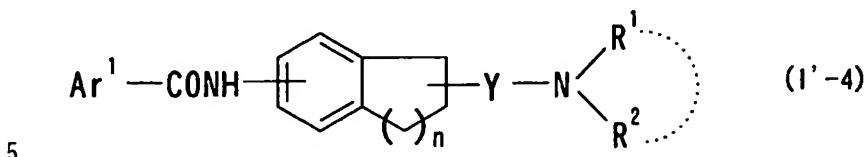


wherein n has the same meaning as defined above, may have further substituents;

provided that N-[2-(N,N-dimethylamino)methyl-6-30 tetralinyl]-4-biphenylcarboxamide is excluded; or a salt

thereof.

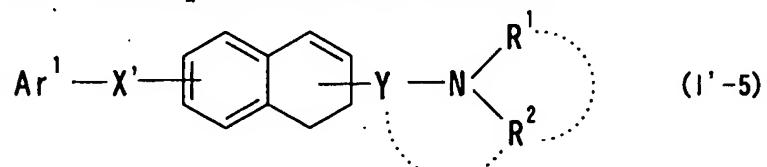
23. A compound according to claim 22, which is of the formula :



wherein R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have
10 substituents; the other symbols have the same meanings as defined in claim 22.

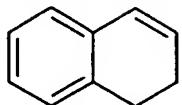
24. A compound according to claim 23, wherein Ar¹ is an aromatic group which may have substituents; and "a
15 hydrocarbon group which may have substituents" for R¹ and R² is "C₁₋₆ alkyl which may have substituents".

25. A compound of the formula :



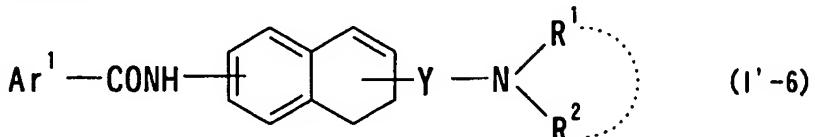
20 wherein Ar¹ is a cyclic group which may have substituents; X' is -CONR^{8c}-, -NR^{8c}CO- or -CH=CH-CONR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl;
Y is a spacer having a main chain of 1 to 6 atoms;
R¹ and R² are independently hydrogen atom or a hydrocarbon
25 group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have
nitrogen-containing hetero ring which may have
30 substituents;

a ring of the formula :



may have further substituents; or a salt thereof.

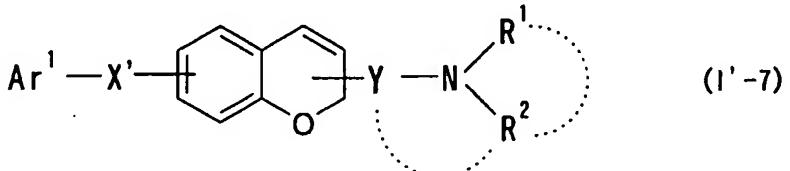
5 26. A compound according to claim 25, which is of the formula :



wherein R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², 10 together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; the other symbols have the same meanings as defined in claim 25.

15 27. A compound according to claim 26, wherein Ar¹ is an aromatic group which may have substituents; and "a hydrocarbon group which may have substituents" for R¹ and R² is "C₁₋₆ alkyl which may have substituents".

20 28. A compound of the formula :

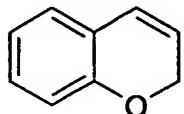


wherein Ar¹ is a cyclic group which may have substituents; X' is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CO-NR^{8c}- or -SO₂NR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl;

25 Y is a spacer having a main chain of 1 to 6 atoms; R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing

hetero ring which may have substituents; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;

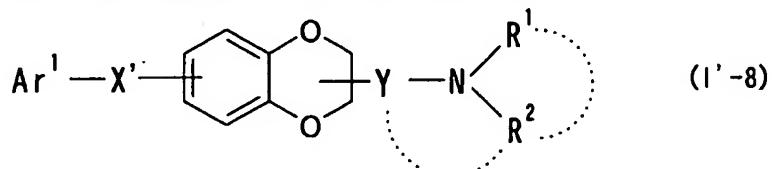
5 a ring of the formula :



may have further substituents;
provided that Ar¹ is not biphenylyl which may be substituted, when X' is -CONH-; or a salt thereof.

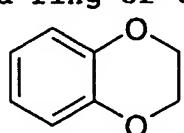
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29. A compound of the formula :



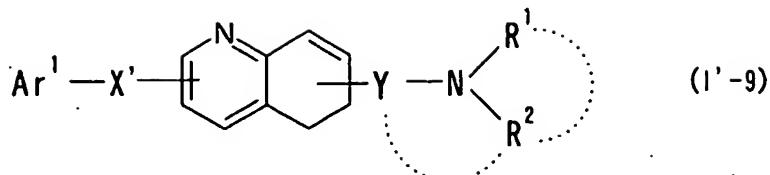
wherein Ar¹ is a cyclic group which may have substituents;
X' is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CONR^{8c}- or -SO₂NR^{8c}- where
15 R^{8c} is hydrogen atom or C₁₋₆ alkyl;
Y is a spacer having a main chain of 1 to 6 atoms;
R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R², together
20 with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;

20 a ring of the formula :



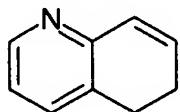
25 may have further substituents; or a salt thereof.

30. A compound of the formula :



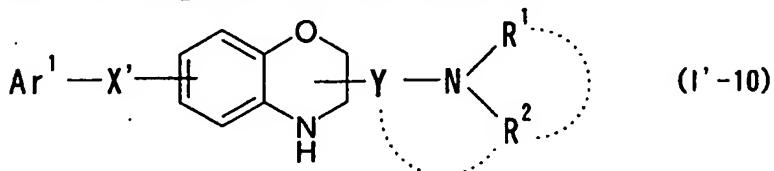
wherein Ar¹ is a cyclic group which may have substituents; X' is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CO-NR^{8c}- or -SO₂NR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl;

5 Y is a spacer having a main chain of 1 to 6 atoms;
 R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R², together
 10 with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents;
 a ring of the formula :



15 may have further substituents; or a salt thereof.

31. A compound of the formula :

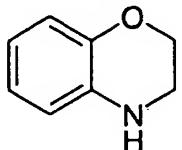


wherein Ar¹ is a cyclic group which may have substituents;

20 X' is -CONR^{8c}-, -NR^{8c}CO-, -CH=CH-CO-NR^{8c}- or -SO₂NR^{8c}- where R^{8c} is hydrogen atom or C₁₋₆ alkyl;
 Y is a spacer having a main chain of 1 to 6 atoms;
 R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with
 25 the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have

substituents;

a ring of the formula :



may have further substituents;

5 provided that Ar¹ is not biphenylyl which may be substituted, when X' is -CONH-; or a salt thereof.

32. A pharmaceutical composition which comprises a compound as defined in any one of claims 18, 19, 22, 25,
10 26, 28, 29, 30 and 31.

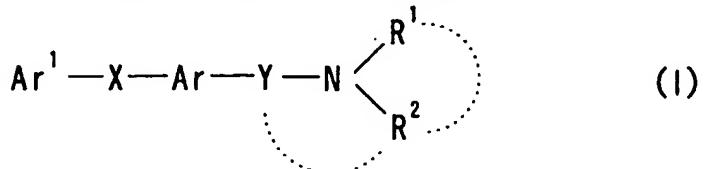
33. A prodrug of a compound as defined in any one of claims 18, 19, 22, 25, 26, 28, 29, 30 and 31.

15 34. A compound according to claim 18, which is N-[2-(N,N-dimethylamino)methyl-6-tetralinyl]-(4'-methoxybiphenyl-4-yl)carboxamide;
4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
20 4'-fluoro-N-[6-(1-piperidinylmethyl)-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]4-carboxamide;
4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
25 (+)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
(-)-4'-fluoro-N-[6-[(N,N-dimethylamino)methyl]-5,6,7,8-tetrahydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
30 4'-chloro-N-[3-[(N,N-dimethylamino)methyl]-2H-chromen-7-yl][1,1'-biphenyl]-4-carboxamide;
4'-fluoro-N-[6-(1-pyrrolidinylmethyl)-7,8-dihydro-2-

naphthalenyl][1,1'-biphenyl]-4-carboxamide;
N-[3-[(dimethylamino)methyl]-2H-chromen-7-yl]-4'-
fluoro[1,1'-biphenyl]-4-carboxamide;
4'-chloro-N-[6-[(dimethylamino)methyl]-5-methyl-7,8-
5 dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide;
6-(4-methoxyphenyl)-N-[5-methyl-6-(1-
pyrrolidinylmethyl)-7,8-dihydro-2-
naphthalenyl]nicotinamide;
4'-chloro-N-[7-[(dimethylamino)methyl]-5,6-dihydro-3-
10 quinolinyl][1,1'-biphenyl]-4-carboxamide;
4-(4-chlorophenyl)-N-[6-(1-pyrrolidinylmethyl)-7,8-
dihydro-2-naphthalenyl]-3,6-dihydro-1(2H)-
pyridinecarboxamide;
N-[6-[(dimethylamino)methyl]-7,8-dihydro-2-
15 naphthalenyl]-4-(4-fluorophenyl)-1-
piperidinecarboxamide;
4-(4-methoxyphenyl)-N-[6-(1-pyrrolidinylmethyl)-5-
methyl-7,8-dihydro-2-naphthalenyl]-1-
piperidinecarboxamide;
20 4'-fluoro-N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-
naphthalenyl][1,1'-biphenyl]-4-carboxamide;
4'-chloro-N-[6-[2-(1-pyrrolidinyl)ethyl]-7,8-dihydro-2-
naphthalenyl][1,1'-biphenyl]-4-carboxamide;
4'-chloro-N-[2-[(dimethylamino)methyl]-3,4-dihydro-2H-
25 1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide;
4-(4-methoxyphenyl)-N-[5-methyl-6-(1-
pyrrolidinylmethyl)-7,8-dihydro-2-naphthalenyl]-1-
piperidinecarboxamide;
4-(4-chlorophenyl)-N-[6-[(4-methyl-1-
30 piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-
piperidinecarboxamide;
4'-chloro-N-[2-[(dimethylamino)methyl]-1H-inden-6-
yl][1,1'-biphenyl]-4-carboxamide;
4'-fluoro-N-[2-(1-pyrrolidinylmethyl)-3,4-dihydro-2H-
35 1,4-benzoxazin-6-yl][1,1'-biphenyl]-4-carboxamide;
4'-fluoro-N-[5-methyl-6-[(4-methyl-1-

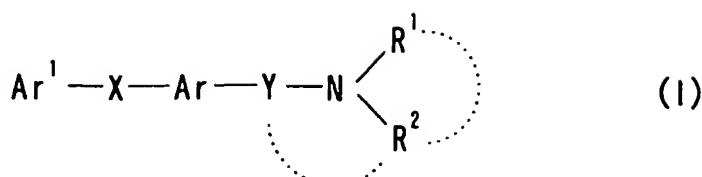
piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide; 4'-chloro-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl][1,1'-biphenyl]-4-carboxamide; or 4-(4-chlorophenyl)-N-[5-methyl-6-[(4-methyl-1-piperazinyl)methyl]-7,8-dihydro-2-naphthalenyl]-1-piperidinecarboxamide.

10 35. A method for preventing or treating diseases caused by a melanin-concentrating hormone in a mammal in need thereof, which comprises administering to said mammal an effective amount of a compound of the formula :



15 wherein Ar¹ is a cyclic group which may have substituents; X is a spacer having a main chain of 1 to 6 atoms; Y is a bond or a spacer having a main chain of 1 to 6 atoms; Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have further
20 substituents;
R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; R² may form a spiro
25 ring together with Ar; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof.

36. A method for preventing or treating obesity in a mammal
30 in need thereof, which comprises administering to said
mammal an effective amount of a compound of the formula :



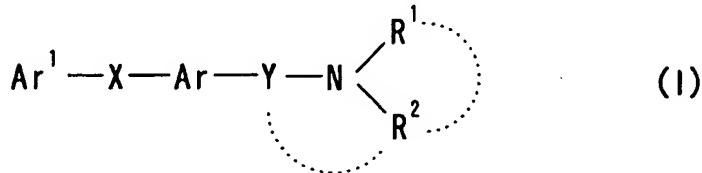
wherein Ar¹ is a cyclic group which may have substituents; X is a spacer having a main chain of 1 to 6 atoms; Y is a bond or a spacer having a main chain of 1 to 6 atoms;

5 Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have further substituents;

10 R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; R² may form a spiro ring together with Ar; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof.

15

37. Use of a compound of the formula :



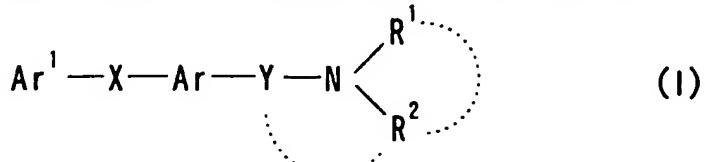
wherein Ar¹ is a cyclic group which may have substituents; X is a spacer having a main chain of 1 to 6 atoms;

20 Y is a bond or a spacer having a main chain of 1 to 6 atoms; Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have further substituents;

25 R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; R² may form a spiro ring together with Ar; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof;

for the manufacture of a pharmaceutical preparation for preventing or treating diseases caused by a melanin-concentrating hormone.

5 38. Use of a compound of the formula :



wherein Ar¹ is a cyclic group which may have substituents; X is a spacer having a main chain of 1 to 6 atoms; Y is a bond or a spacer having a main chain of 1 to 6 atoms; 10 Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have further substituents; R¹ and R² are independently hydrogen atom or a hydrocarbon group which may have substituents; R¹ and R², together with 15 the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; R² may form a spiro ring together with Ar; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof; 20 for the manufacture of a pharmaceutical preparation for preventing or treating obesity.

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Matsushiro 3-chome, Tsukuba-shi, Ibaraki 305-0035
(JP). TAKEKAWA, Shiro [JP/JP]; 5-3-B305, Umezono
2-chome, Tsukuba-shi, Ibaraki 305-0045 (JP). ISHI-
HARA, Yuji [JP/JP]; 12-30-305, Ninomiya 1-chome,
Tsukuba-shi, Ibaraki 305-0051 (JP).

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(74) Agents: TAKAHASHI, Shuichi et al.; Osaka Plant of
Takeda Chemical Industries, Ltd., 17-85, Jusohonmachi
2-chome, Yodogawa-ku, Osaka-shi, Osaka 532-0024 (JP).

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IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG,
CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

(71) Applicant (*for all designated States except US*): TAKEDA
CHEMICAL INDUSTRIES, LTD. [JP/JP]; 1-1,
Doshomashi 4-chome, Chuo-ku, Osaka-shi, Osaka
541-0045 (JP).

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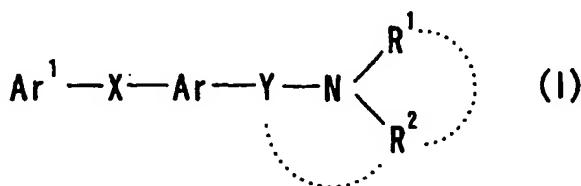
(72) Inventors; and
(75) Inventors/Applicants (*for US only*): KATO, Kaneyoshi
[JP/JP]; 2-40, Maruyamadai 2-chome, Kawanishi-shi,
Hyogo 666-0152 (JP). TERAUCHI, Jun [JP/JP]; 3-5-204,
Hachizuka 3-chome, Ikeda-shi, Osaka 563-0024 (JP).
MORI, Masaaki [JP/JP]; 7-9-702, Kasuga 1-chome,
Tsukuba-shi, Ibaraki 305-0821 (JP). SUZUKI, Nobuhiro
[JP/JP]; 1077-50, Oaza-yatabe, Tsukuba-shi, Ibaraki
305-0861 (JP). SHIMOMURA, Yukio [JP/JP]; 12-1-410,

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which may have substituents; R¹ and R², together with the adjacent nitrogen atom, may form a nitrogen-containing hetero ring which may have substituents; R² may form a spiro ring together with Ar; or R², together with the adjacent nitrogen atom and Y, may form a nitrogen-containing hetero ring which may have substituents; or a salt thereof; which is useful as an agent for preventing or treating obesity, etc.

(57) Abstract: A melanin-concentrating hormone antagonist which comprises a compound of formula (I) wherein Ar¹ is a cyclic group which may have substituents; X is a spacer having a main chain of 1 to 6 atoms; Y is a bond or a spacer having a main chain of 1 to 6 atoms; Ar is a monocyclic aromatic ring which may be condensed with a 4 to 8 membered non-aromatic ring, and may have further substituents; R¹ and R² are independently hydrogen atom or a hydrocarbon group

INTERNATIONAL SEARCH REPORT

International Application No
PCT/JP 00/06375

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|-------------------------------------|------------|------------|------------|------------|------------|
| A. CLASSIFICATION OF SUBJECT MATTER | | | | | |
| IPC 7 | C07C235/42 | C07C235/84 | C07D209/48 | C07C237/40 | C07C275/42 |
| | C07C233/44 | A61K31/16 | A61K31/40 | | |

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 C07C C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

CHEM ABS Data, WPI Data, EPO-Internal, BEILSTEIN Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

| Category | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
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| X | WO 95 32967 A (SMITHKLINE BEECHAM PLC ;HAM PETER (GB); GASTER LARAMIE MARY (GB);) 7 December 1995 (1995-12-07) cited in the application claims; examples --- | 1-5,7, 13-16 |
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Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

& document member of the same patent family

Date of the actual completion of the international search

19 December 2000

Date of mailing of the international search report

15.01.01

Name and mailing address of the ISA

European Patent Office, P.O. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel: (+31-70) 340-2040, Tx. 31 651 epo nl.
Fax: (+31-70) 340-3016

Authorized officer

Seufert, G

INTERNATIONAL SEARCH REPORT

| |
|---|
| International Application No PCT/JP 00/06375 |
|---|

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

| Category | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|----------|---|--------------------------------|
| X | DE 21 08 185 A (TROPONWERKE DINKLAGE & CO.) 7 September 1972 (1972-09-07) claims; examples --- | 1-4, 8-16, 18, 28, 32 |
| X | DE 24 48 257 A (TROPONWERKE DINKLAGE & CO) 22 April 1976 (1976-04-22) claims; examples --- | 1-4, 8-16 |
| X | DE 25 02 588 A (TROPONWERKE DINKLAGE & CO) 29 July 1976 (1976-07-29) claims; examples --- | 1-4, 8-16 |
| X | R. E. MEWSHAW ET AL.: "New Generation Dopaminergic Agents. 1. Discovery of a Novel Scaffold Which Embraces the D2 Agonist Pharmacophore. Structure-Activity Relationship of a Series of 2-(Aminomethyl)chromans" J. MED. CHEM., vol. 40, no. 26, 1997, pages 4235-56, XP002155829 * page 4248, right column, example 22b - page 4249, left column, example 39a * --- | 1-4, 7, 9, 11-13 |
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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

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| X | EP 0 920 864 A (PFIZER PROD INC) 9 June 1999 (1999-06-09) * page 3, line 10-14, 30-36; claims; examples * ---- | 1,4,7, 13-17, 35-38 |
| A | QU D ET AL: "A ROLE FOR MELANIN-CONCENTRATING HORMONE IN THE CENTRAL REGULATION OF FEEDING BEHAVIOUR" NATURE, GB, MACMILLAN JOURNALS LTD. LONDON, vol. 380, 21 March 1996 (1996-03-21), pages 243-247, XP002037981 ISSN: 0028-0836 the whole document ---- | 1-38 |

INTERNATIONAL SEARCH REPORTnational application No.
PCT/JP 00/06375**Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)**

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
Although claims 35 and 36 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. Claims Nos.: 1-33(partly), 35-38(partly)
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
see FURTHER INFORMATION sheet PCT/ISA/210
3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/SA/ 210

Continuation of Box I.2

Claims Nos.: 1-33(partly), 35-38(partly)

The initial phase of the search revealed a very large number of documents relevant to the issue of novelty (attention is drawn to the fact that claim 1 as it is drafted has to be considered purely as a compound claim). So many documents were retrieved (a few have been cited as a mere random selection) that it is impossible to determine which parts of the claim(s) may be said to define subject-matter for which protection might legitimately be sought (Article 6 PCT). For these reasons, a meaningful search over the whole breadth of the claim(s) is impossible. Consequently, the search has been restricted to:

compounds to formula I' (see claim 18) with X = C(O)N, Y= -(CH₂)₂- whereby the substituents A₁-X- and YNR₁R₂ may not be attached in alpha-position to the C-atoms shared by the condensed ring (Ar'). Ar', Ar and R₁/R₂ are as defined in claim 18.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

INTERNATIONAL SEARCH REPORT

 International Application No
 PCT/JP 00/06375

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